



Submission to
enter the
national stage
under 35 U.S.C.
§ 371.

JC17 Rec'd PCT/PTO 04 MAY 2001

FORM PTO-1390
(REV. 11-2000)

U S DEPARTMENT OF COMMERCE PATENT AND TRADEMARK
26959
PATENT & TRADEMARK OFFICE

TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371

ATTORNEY'S DOCKET NUMBER
CRT-5463 (1417S P 585)

U S APPLICATION NO (If known, see 37 CFR 1.5)

09/831121

INTERNATIONAL APPLICATION NO.
PCT/US99/25955

INTERNATIONAL FILING DATE
4 November 1999

PRIORITY DATE CLAIMED
4 November 1998

TITLE OF INVENTION

ELEMENT PROVIDED WITH A FIBRIN LAYER, PREPARATION AND USE THEREOF

APPLICANT(S) FOR DO/EO/US

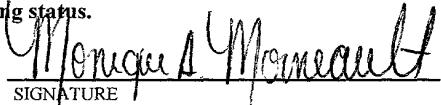
Yves Delmotte, Nathalie Belot, Pierre Vermeulen and Nicole Tasiaux

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (21) indicated below.
4. The US has been elected by the expiration of 19 months from the priority date (Article 31).
5. A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. is attached hereto (required only if not communicated by the International Bureau).
 - b. has been communicated by the International Bureau.
 - c. is not required, as the application was filed in the United States Receiving Office (RO/US).
6. An English language translation of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. is attached hereto.
 - b. has been previously submitted under 35 U.S.C. 154(d)(4).
7. Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. are attached hereto (required only if not communicated by the International Bureau).
 - b. have been communicated by the International Bureau.
 - c. have not been made; however, the time limit for making such amendments has NOT expired.
 - d. have not been made and will not be made.
8. An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11 to 20 below concern document(s) or information included:

11. An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. A **FIRST** preliminary amendment.
14. A **SECOND** or **SUBSEQUENT** preliminary amendment.
15. A substitute specification.
16. A change of power of attorney and/or address letter.
17. A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.
18. A second copy of the published international application under 35 U.S.C. 154(d)(4).
19. A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).
20. Other items or information:
 - Notification of Transmittal of the International Preliminary Examination Report and International Preliminary Examination Report
 - International Search Report

U.S. APPLICATION NO. (Unknown see 37 CFR 1.5)	INTERNATIONAL APPLICATION NO PCT/US99/25955	ATTORNEY'S DOCKET NUMBER CRT-5463(1417S P 585)																										
21. <input checked="" type="checkbox"/> The following fees are submitted:		CALCULATIONS PTO USE ONLY																										
BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)):																												
Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO. \$1000.00																												
<input checked="" type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$860.00																												
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International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$690.00																												
International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) \$100.00																												
ENTER APPROPRIATE BASIC FEE AMOUNT =																												
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input checked="" type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).		\$ 130																										
<table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th>CLAIMS</th> <th>NUMBER FILED</th> <th>NUMBER EXTRA</th> <th>RATE</th> <th>\$</th> </tr> </thead> <tbody> <tr> <td>Total claims</td> <td>89 - 20 =</td> <td>69</td> <td>x \$18.00</td> <td>\$ 1,242</td> </tr> <tr> <td>Independent claims</td> <td>6 - 3 =</td> <td>3</td> <td>x \$80.00</td> <td>\$ 240</td> </tr> <tr> <td colspan="4">MULTIPLE DEPENDENT CLAIM(S) (if applicable)</td> <td>+ \$270.00</td> </tr> <tr> <td colspan="4" style="text-align: center;">TOTAL OF ABOVE CALCULATIONS</td> <td style="text-align: right;">\$ 2,472</td> </tr> </tbody> </table>				CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	\$	Total claims	89 - 20 =	69	x \$18.00	\$ 1,242	Independent claims	6 - 3 =	3	x \$80.00	\$ 240	MULTIPLE DEPENDENT CLAIM(S) (if applicable)				+ \$270.00	TOTAL OF ABOVE CALCULATIONS				\$ 2,472
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<input type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.		\$																										
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Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).		\$																										
		TOTAL NATIONAL FEE = \$ 2,472																										
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property		\$																										
		TOTAL FEES ENCLOSED = \$ 2,472																										
		Amount to be refunded:	\$																									
		charged:	\$																									
<p>a. <input checked="" type="checkbox"/> A check in the amount of \$ <u>2,472.00</u> to cover the above fees is enclosed.</p> <p>b. <input type="checkbox"/> Please charge my Deposit Account No. _____ in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed.</p> <p>c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>23-0280</u>. A duplicate copy of this sheet is enclosed.</p> <p>d. <input type="checkbox"/> Fees are to be charged to a credit card. WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.</p>																												
<p>NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137 (a) or (b)) must be filed and granted to restore the application to pending status.</p> <p>SEND ALL CORRESPONDENCE TO: Janice Guthrie, Ph.D. (Reg. No. 35,170) BAXTER HEALTHCARE CORPORATION P.O. Box 15210 Irvine, CA 92623-5210 Telephone: 949.250.6828 Facsimile: 949.250.6850</p>																												
 SIGNATURE Monique A. Morneault NAME 37,893 REGISTRATION NUMBER																												

Attorney Docket No. CRT-5463 (1417S P 585)
/121835

JC03 Rec'd PCT/PTC

04 MAY 2001
PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In Re U.S. Patent Application of:)

Yves Delmotte, Nathalie Belot, Pierre Vermeulen and Nicole Tasiaux)

From PCT/US99/25955 (Filed 4 November 1999))

And Belgian Application No. BE 9800796 (Filed 4 November 1998))

For: Element Provided With a Fibrin Layer, Preparation and Use Thereof)

Mailed: May 4, 2001)

PRELIMINARY AMENDMENT

BOX PCT
Commissioner for Patents
Washington, DC 20231

Dear Commissioner:

Before examination of this Application, please make the following amendments.

IN THE SPECIFICATION:

Amend the specification by inserting before the first line the sentence:

-- This Application is a U.S. National §371 filing from Belgian Application No. BE 9800796, filed November 4, 1998, now pending (which is hereby incorporated by reference). --

On Page 1, after the title, please add the heading -- **DESCRIPTION** --.

On Page 1, after the above new heading is added, please add the heading -- **Technical Field** --.

On Page 1, after line 12, please add the heading -- **Background of the Invention** --.

On Page 2, after line 2, please add the heading -- **Summary of the Invention** --.

On Page 18, after line 13, please add the heading -- **Brief Description of the Drawings** --.

On Page 19, after line 19, please add the heading -- **Detailed Description of the Preferred Embodiments** --.

IN THE CLAIMS:

Please cancel claims 1-45 and add the claims 46-134:

46. An element comprising:
a support having a face;
the face having at least two pores spaced from one another to define a node spacing; and
a fibrin network substantially free of unbound fibrinogen covering a portion of the face.
47. The element according to claim 46 wherein the support is substantially hydrophobic.
48. The element according to claim 46 wherein the face has a thickness from about 0.1 mm to about 5 mm.
49. The element according to claim 46 wherein the fibrin network is positioned over a portion of the pores.
50. The element of claim 46 wherein the pores extend through the thickness of the support.
51. The element according to claim 46 wherein the node spacing is from about 5 μm to about 100 μm .
52. The element of claim 46 wherein the fibrin network is substantially uniform and homogeneous.
53. The element according to claim 52 wherein the fibrin network contains less than 1% by weight of fibrinogen that has not reacted.
54. The element according to claim 53 wherein the fibrin network contains less than 0.5% by weight of fibrinogen that has not reacted.

55. The element according to claim 54 wherein the fibrin network contains less than 0.1% by weight of fibrinogen that has not reacted.

56. The element according to claim 55 wherein the fibrin network extends through the pores to a depth of the support of at least 2 μm .

57. The element according to claim 56 wherein the node spacing is substantially homogeneous and uniform.

58. The element according to claim 46 wherein the fibrinogen extends through the thickness of the fibrin network to a depth of at least 10 μm .

59. The element according to claim 58 wherein the fibrinogen extends through the thickness of the fibrin network to a depth of at least 20 μm .

60. The element according to claim 46 wherein the fibrin network further includes a first face and a second face.

61. The element according to claim 60 wherein first face is in contact with the support.

62. The element according to claim 61 wherein second face is stabilized by at least partial cross-linking to form a network of adjacent alveoli.

63. The element according to claim 60 wherein the fibrin network is provided with cells.

64. The element according to claim 63 wherein the cells mediate cell-fibrin bonds.

65. The element according to claim 60 wherein the fibrin network is provided with

proteins.

66. The element according to claim 65 wherein the proteins mediate cell-fibrin bonds.
67. The element according to claim 62 wherein the at least partially cross-linked fibrin network is 0.5 to 100 μm thick when measured in the dry state.
68. The element according to claim 67 wherein the at least partially cross-linked fibrin network is 2.5 to 50 μm thick when measured in the dry state.
69. The element according to claim 62 wherein the alveoli are formed between the cross-linked fibrin.
70. The element according to claim 69 wherein the alveoli have a volume from about 5 μm^3 to about 25 μm^3 .
71. The element according to claim 70 wherein the alveoli have an average thickness from about 1 μm to about 5 μm when measured in the dry state.
72. The element according to claim 46 wherein the pores further include inner faces.
73. The element according to claim 72 wherein the inner faces are at least partially covered by a water-soluble or substantially water-soluble protein.
74. The element according to claim 46 wherein the fibrin network is at least partially covered by a water-soluble or substantially water-soluble protein.
75. The element according to claim 72 wherein the pores are covered by a water-soluble or miscible polar organic additive.

76. The element according to claim 62 wherein the cross-linked fibrin is at least partially covered by a water-soluble or miscible polar additive.

77. The element according to claim 76 wherein the water-soluble or miscible polar additive is selected in the group comprising glycerol, sugars and mixtures thereof.

78. The element according to claim 46 wherein the fibrin network has a moisture content of less than 0.5%.

79. The element according to claim 46 wherein the fibrin network contains fibronectin.

80. The element according to claim 79 wherein the fibronectin content is from about 0.5% to about 15%.

81. The element according to claim 46 wherein the fibrin network contains calcium.

82. The element according to claim 81 wherein the calcium content is from about 1 μ g to about 100 μ g per cm^3 of the fibrin network volume.

83. The element according to claim 81 wherein the calcium is calcium chloride.

84. The element according to claim 46 wherein the support further includes a second fibrin network superimposed on a first fibrin network.

85. The element according to claim 84 wherein the first fibrin network is in contact with the support.

86. The element according to claim 85 wherein the first fibrin network contains alveoli.

87. The element according to claim 84 wherein the second fibrin network contains alveoli.

88. The element according to claim 84 wherein the alveoli of the first fibrin network have larger volumes than the alveoli of the second fibrin network.

89. The element according to claim 46 wherein the support is biocompatible and/or biodegradable.

90. A method for producing an element, the method comprising placing a first porous portion of a support in contact with an aqueous solution and submitting a second porous portion of the support to a substantially uniform and homogenous suction force.

91. The method according to claim 90 wherein the second porous portion is positioned opposite to the first porous portion.

92. The method according to claim 90 wherein the aqueous solution contains fibrin.

93. The method according to claim 92 wherein the aqueous solution contains fibrinogen.

94. The method according to claim 93 wherein the aqueous solution contains water.

95. The method according to claim 90 wherein the suction force pulls the water through the first porous portion.

96. The method according to claim 95 wherein the suction force deposits a fibrin network on the first porous portion.

97. The method according to claim 96 wherein the fibrin network is deposited

substantially uniformly and homogeneously.

98. The method according to claim 97 wherein the fibrin penetrates the first porous portion.

99. The method according to claim 90 wherein the second porous portion is submitted to a pressure of less than about 0.8×10^5 Pa.

100. The method according to claim 99 wherein the second porous portion is submitted to a pressure of less than about 0.5×10^5 Pa.

101. The method according to claim 90 wherein a pressure difference is created between the first porous portion and the second porous portion.

102. The method according to claim 101 wherein a pressure difference is at least 0.3×10^5 Pa.

103. The method according to claim 101 wherein the second porous portion is intermittently submitted to a first pressure and a second pressure.

104. The method according to claim 103 wherein the first pressure is less than about 0.8×10^5 Pa.

105. The method according to claim 103 wherein the second pressure is less than about 0.8×10^5 Pa.

106. The method according to claim 103 wherein the first pressure is at least 5% higher than the second pressure.

107. The method according to claim 101 wherein the second porous portion is exposed to a temperature from about 0°C to about 100°C.

108. The method according to claim 107 wherein the second porous portion is exposed to a temperature from about 15°C to about 60°C.

109. The method according to claim 90 wherein the second porous portion is submitted to reverse osmosis.

110. The method according to claim 109 wherein the reverse osmosis causes the diffusion of at least the solution water through the first porous portion.

111. The method according to claim 90 wherein the solution contains from about 5 to about 20 mg/ml of fibrinogen-containing materials.

112. The method according to claim 111 wherein the solution contains from about 0.01 to about 10 units of thrombin per ml.

113. The method according to claim 112 wherein the solution contains from about 0.1 to about 10 units of factor XIII per ml.

114. The method according to claim 113 wherein the solution contains from about 1 to about 40 millimoles of calcium chloride per ml.

115. The method according to claim 114 wherein the solution contains from about 0 % to about 20% by weight of a water-soluble or miscible polar organic additive.

116. The method according to claim 115 wherein the solution contains from about 5% to about 10% by weight of a water-soluble or miscible polar organic additive.

117. The method according to claim 116 wherein the additive is glycerol.

118. The method according to claim 90 further comprising:

a first step wherein the first porous portion is placed in contact with a solution containing fibrin and/or fibrinogen-containing materials;

a second step wherein the second porous portion is submitted to a suction force to diffuse at least the solution water through the thickness of the porous support and homogeneously and uniformly having penetration of fibrin or fibrinogen across the thickness of the porous support; and

a third step wherein the fibrin and/or fibrinogen network is stabilized by at least partial cross-linking to form a network of adjacent alveoli.

119. The method of claim 118 wherein the solution is in motion.

120. The method of claim 119 wherein the solution contains a wetting agent.

121. The method of claim 118 wherein the fibrin network is submitted to a washing step.

122. The method of claim 121 wherein the fibrin network is submitted to a drying step.

123. The method of claim 122 wherein the drying step is accomplished at least partially by lyophilization.

124. The method of claim 123 wherein the drying step is accomplished at a temperature from about -30°C to about -100°C.

125. The method of claim 124 wherein the drying step is accomplished at a temperature from about -40°C to about -70°C.

126. The method of claim 118 wherein the fibrin or fibrinogen-containing materials have a

controlled concentration.

127. The method of claim 126 wherein the concentration ensures a substantially constant water diffusion through the support.

128. The method of claim 90 wherein the support is biocompatible and/or biodegradable.

129. The method of claim 90 wherein the porous portion is treated with an aqueous solution before contacting the fibrin and/or fibrinogen-containing materials.

130. The method of claim 129 wherein the solution contains a wetting agent, a protein or a polar organic additive, or a mixture thereof.

131. A filter including a membrane element comprising:
a support having a face;
the face having at least two pores spaced from one another to define a node spacing; and
a fibrin network substantially free of unbound fibrinogen covering a portion of the face.

132. A bioreactor including a membrane element comprising:
a support having a face;
the face having at least two pores spaced from one another to define a node spacing; and
a fibrin network substantially free of unbound fibrinogen covering a portion of the face.

133. An implant consisting of an element comprising:
a support having a face;
the face having at least two pores spaced from one another to define a node spacing; and
a fibrin network substantially free of unbound fibrinogen covering a portion of the face.

134. An artificial skin produced from an element comprising:

a support having a face;

the face having at least two pores spaced from one another to define a node spacing; and
a fibrin network substantially free of unbound fibrinogen covering a portion of the face.

REMARKS

Applicants respectfully request that this Amendment be entered and examined. The Headings have been added to conform the Application to United States format. The Claims have also been amended to conform to United States practice. No new matter has been introduced by these Amendments.

The Commissioner is hereby authorized to charge payment of any deficit in fees associated with this Amendment to Deposit Account No. 23-0280. A duplicate copy of this sheet is enclosed for that purpose.

Please address all future correspondence to the undersigned attorney and domestic representative for Applicant.

Respectfully submitted,

Date: May 4, 2001

By: Monique A. Morneau
Joseph A. Fuchs, Registration No. 34,604
Monique A. Morneau, Registration No. 37,893
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Chicago, Illinois 60606-6630
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Attorney for Applicants

Express Mail Label No. EL619829524US

Date of Deposit: May 4, 2001

I hereby certify that this paper or fee is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 C.F.R. § 1.10, postage prepaid, on the date indicated above and is addressed to: Box PCT, Commissioner for Patents, Washington, DC 20231.

Julie M. Weisenberger.
Julie M. Weisenberger

Attorney Docket No. CRT-5463 (1417S P 585)
/121835

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IN THE CLAIMS:

Please cancel claims 1-45 and add the claims 46-134:

46. An element comprising:
a support having a face;
the face having at least two pores spaced from one another to define a node spacing; and
a fibrin network substantially free of unbound fibrinogen covering a portion of the face.
47. The element according to claim 46 wherein the support is substantially hydrophobic.
48. The element according to claim 46 wherein the face has a thickness from about 0.1 mm to about 5 mm.
49. The element according to claim 46 wherein the fibrin network is positioned over a portion of the pores.
50. The element of claim 46 wherein the pores extend through the thickness of the support.
51. The element according to claim 46 wherein the node spacing is from about 5 μm to about 100 μm .
52. The element of claim 46 wherein the fibrin network is substantially uniform and homogeneous.
53. The element according to claim 52 wherein the fibrin network contains less than 1% by weight of fibrinogen that has not reacted.
54. The element according to claim 53 wherein the fibrin network contains less than 0.5% by weight of fibrinogen that has not reacted.

55. The element according to claim 54 wherein the fibrin network contains less than 0.1% by weight of fibrinogen that has not reacted.

56. The element according to claim 55 wherein the fibrin network extends through the pores to a depth of the support of at least 2 μm .

57. The element according to claim 56 wherein the node spacing is substantially homogeneous and uniform.

58. The element according to claim 46 wherein the fibrinogen extends through the thickness of the fibrin network to a depth of at least 10 μm .

59. The element according to claim 58 wherein the fibrinogen extends through the thickness of the fibrin network to a depth of at least 20 μm .

60. The element according to claim 46 wherein the fibrin network further includes a first face and a second face.

61. The element according to claim 60 wherein first face is in contact with the support.

62. The element according to claim 61 wherein second face is stabilized by at least partial cross-linking to form a network of adjacent alveoli.

63. The element according to claim 60 wherein the fibrin network is provided with cells.

64. The element according to claim 63 wherein the cells mediate cell-fibrin bonds.

65. The element according to claim 60 wherein the fibrin network is provided with

proteins.

66. The element according to claim 65 wherein the proteins mediate cell-fibrin bonds.
67. The element according to claim 62 wherein the at least partially cross-linked fibrin network is 0.5 to 100 μm thick when measured in the dry state.
68. The element according to claim 67 wherein the at least partially cross-linked fibrin network is 2.5 to 50 μm thick when measured in the dry state.
69. The element according to claim 62 wherein the alveoli are formed between the cross-linked fibrin.
70. The element according to claim 69 wherein the alveoli have a volume from about 5 μm^3 to about 25 μm^3 .
71. The element according to claim 70 wherein the alveoli have an average thickness from about 1 μm to about 5 μm when measured in the dry state.
72. The element according to claim 46 wherein the pores further include inner faces.
73. The element according to claim 72 wherein the inner faces are at least partially covered by a water-soluble or substantially water-soluble protein.
74. The element according to claim 46 wherein the fibrin network is at least partially covered by a water-soluble or substantially water-soluble protein.
75. The element according to claim 72 wherein the pores are covered by a water-soluble or miscible polar organic additive.

76. The element according to claim 62 wherein the cross-linked fibrin is at least partially covered by a water-soluble or miscible polar additive.

77. The element according to claim 76 wherein the water-soluble or miscible polar additive is selected in the group comprising glycerol, sugars and mixtures thereof.

78. The element according to claim 46 wherein the fibrin network has a moisture content of less than 0.5%.

79. The element according to claim 46 wherein the fibrin network contains fibronectin.

80. The element according to claim 79 wherein the fibronectin content is from about 0.5% to about 15%.

81. The element according to claim 46 wherein the fibrin network contains calcium.

82. The element according to claim 81 wherein the calcium content is from about 1 μ g to about 100 μ g per cm^3 of the fibrin network volume.

83. The element according to claim 81 wherein the calcium is calcium chloride.

84. The element according to claim 46 wherein the support further includes a second fibrin network superimposed on a first fibrin network.

85. The element according to claim 84 wherein the first fibrin network is in contact with the support.

86. The element according to claim 85 wherein the first fibrin network contains alveoli.

87. The element according to claim 84 wherein the second fibrin network contains alveoli.

88. The element according to claim 84 wherein the alveoli of the first fibrin network have larger volumes than the alveoli of the second fibrin network.

89. The element according to claim 46 wherein the support is biocompatible and/or biodegradable.

90. A method for producing an element, the method comprising placing a first porous portion of a support in contact with an aqueous solution and submitting a second porous portion of the support to a substantially uniform and homogenous suction force.

91. The method according to claim 90 wherein the second porous portion is positioned opposite to the first porous portion.

92. The method according to claim 90 wherein the aqueous solution contains fibrin.

93. The method according to claim 92 wherein the aqueous solution contains fibrinogen.

94. The method according to claim 93 wherein the aqueous solution contains water.

95. The method according to claim 90 wherein the suction force pulls the water through the first porous portion.

96. The method according to claim 95 wherein the suction force deposits a fibrin network on the first porous portion.

97. The method according to claim 96 wherein the fibrin network is deposited

substantially uniformly and homogeneously.

98. The method according to claim 97 wherein the fibrin penetrates the first porous portion.

99. The method according to claim 90 wherein the second porous portion is submitted to a pressure of less than about 0.8×10^5 Pa.

100. The method according to claim 99 wherein the second porous portion is submitted to a pressure of less than about 0.5×10^5 Pa.

101. The method according to claim 90 wherein a pressure difference is created between the first porous portion and the second porous portion.

102. The method according to claim 101 wherein a pressure difference is at least 0.3×10^5 Pa.

103. The method according to claim 101 wherein the second porous portion is intermittently submitted to a first pressure and a second pressure.

104. The method according to claim 103 wherein the first pressure is less than about 0.8×10^5 Pa.

105. The method according to claim 103 wherein the second pressure is less than about 0.8×10^5 Pa.

106. The method according to claim 103 wherein the first pressure is at least 5% higher than the second pressure.

107. The method according to claim 101 wherein the second porous portion is exposed to a temperature from about 0°C to about 100°C.

108. The method according to claim 107 wherein the second porous portion is exposed to a temperature from about 15°C to about 60°C.

109. The method according to claim 90 wherein the second porous portion is submitted to reverse osmosis.

110. The method according to claim 109 wherein the reverse osmosis causes the diffusion of at least the solution water through the first porous portion.

111. The method according to claim 90 wherein the solution contains from about 5 to about 20 mg/ml of fibrinogen-containing materials.

112. The method according to claim 111 wherein the solution contains from about 0.01 to about 10 units of thrombin per ml.

113. The method according to claim 112 wherein the solution contains from about 0.1 to about 10 units of factor XIII per ml.

114. The method according to claim 113 wherein the solution contains from about 1 to about 40 millimoles of calcium chloride per ml.

115. The method according to claim 114 wherein the solution contains from about 0 % to about 20% by weight of a water-soluble or miscible polar organic additive.

116. The method according to claim 115 wherein the solution contains from about 5% to about 10% by weight of a water-soluble or miscible polar organic additive.

117. The method according to claim 116 wherein the additive is glycerol.

118. The method according to claim 90 further comprising:

a first step wherein the first porous portion is placed in contact with a solution containing fibrin and/or fibrinogen-containing materials;

a second step wherein the second porous portion is submitted to a suction force to diffuse at least the solution water through the thickness of the porous support and homogeneously and uniformly having penetration of fibrin or fibrinogen across the thickness of the porous support; and

a third step wherein the fibrin and/or fibrinogen network is stabilized by at least partial cross-linking to form a network of adjacent alveoli.

119. The method of claim 118 wherein the solution is in motion.

120. The method of claim 119 wherein the solution contains a wetting agent.

121. The method of claim 118 wherein the fibrin network is submitted to a washing step.

122. The method of claim 121 wherein the fibrin network is submitted to a drying step.

123. The method of claim 122 wherein the drying step is accomplished at least partially by lyophilization.

124. The method of claim 123 wherein the drying step is accomplished at a temperature from about -30°C to about -100°C.

125. The method of claim 124 wherein the drying step is accomplished at a temperature from about -40°C to about -70°C.

126. The method of claim 118 wherein the fibrin or fibrinogen-containing materials have a

controlled concentration.

127. The method of claim 126 wherein the concentration ensures a substantially constant water diffusion through the support.

128. The method of claim 90 wherein the support is biocompatible and/or biodegradable.

129. The method of claim 90 wherein the porous portion is treated with an aqueous solution before contacting the fibrin and/or fibrinogen-containing materials.

130. The method of claim 129 wherein the solution contains a wetting agent, a protein or a polar organic additive, or a mixture thereof.

131. A filter including a membrane element comprising:

a support having a face;

the face having at least two pores spaced from one another to define a node spacing; and a fibrin network substantially free of unbound fibrinogen covering a portion of the face.

132. A bioreactor including a membrane element comprising:

a support having a face;

the face having at least two pores spaced from one another to define a node spacing; and a fibrin network substantially free of unbound fibrinogen covering a portion of the face.

133. An implant consisting of an element comprising:

a support having a face;

the face having at least two pores spaced from one another to define a node spacing; and a fibrin network substantially free of unbound fibrinogen covering a portion of the face.

134. An artificial skin produced from an element comprising:

a support having a face;

the face having at least two pores spaced from one another to define a node spacing; and
a fibrin network substantially free of unbound fibrinogen covering a portion of the face.

REMARKS

Applicants respectfully request that this Amendment be entered and examined. The Headings have been added to conform the Application to United States format. The Claims have also been amended to conform to United States practice. No new matter has been introduced by these Amendments.

The Commissioner is hereby authorized to charge payment of any deficit in fees associated with this Amendment to Deposit Account No. 23-0280. A duplicate copy of this sheet is enclosed for that purpose.

Please address all future correspondence to the undersigned attorney and domestic representative for Applicant.

Respectfully submitted,

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Julie M. Weisenberger.
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Element provided with a fibrin layer,
preparation and use thereof.

The invention relates to an element having
5 a fibrin-based layer, said element comprising (a) a hydrophobic or substantially hydrophobic support which has a porous part with a thickness of 0.1 to 5 mm, and whose pores, extending across its thickness, have a node spacing of 5 to 100 μ m, one face of said porous part
10 being treated with a fibrin and/or fibrinogen-based compound, and (b) a fibrin-based layer covering said treated face of the support.

Such elements are known from WO96/07444 and from US 5298255. These known elements are prepared by simply immersing a support in a solution containing fibrinogen and thrombin or by pushing such solution through a porous support. These known elements, when prepared by simple immersion, have a substantially compact fibrin layer and have little or no fibrin in the support pores, or have fibrin in the pores having greater diameters and substantially no fibrin in the pores having smaller diameters, due to an easier passage of fibrin through the pores with greater diameters. This easier passage of fibrin through the pores with greater diameters causes a lack of homogeneity and/or uniformity.

Such lack of homogeneity or uniformity with respect to the presence of fibrin in the support

pores has proved, in some cases, to affect cell attachment.

This invention aims at obviating these drawbacks and essentially relates to an element as described in the first paragraph of this specification, said element being characterized in that said fibrin-based layer is substantially uniform and homogeneous on said treated surface. The fibrin layer according to the present invention is characterized by the lack of fibrinogen, unbound from the fibrin layer. The lack of fibrinogen on the fibrin layer may be detected by the absence of the γ band in the electrophoresis diagram. The lack of fibrinogen in the fibrin layer is caused by the fact that any fibrinogen which has not reacted to form the fibrin layer is sucked through the porous support. Hence, the element according to the invention is characterized in that, at the contact surface between the fibrin layer and the support, there is substantially preferably no fibrinogen which has not reacted. For example, the fibrin layer of the element according to the invention contains less than 2% by weight of fibrinogen which has not reacted to form a fibrin network, preferably less than 1%, particularly less than 0.5% and more particularly less than 0,1%.

Advantageously, the fibrin layer and at least a support layer extending across a thickness of 10 μ m contains less than 1% by weight, advantageously less than 0.5%, preferably less than 0.1% by weight of fibrinogen which has not reacted, with respect to the weight of the fibrin layer. Preferably, fibrin extends across the thickness of the treated porous part of the support, from said treated face to a depth of at least 2 μ m, both through the pores having an average diameter of 10 to 20 μ m and through the pores having an average diameter of more than 20 μ m.

In accordance with a particular embodiment, in which the porous part of the support has

a substantially homogeneous and uniform porosity over the treated face, some fibrin extends homogeneously and uniformly across the thickness of the porous part of the support to a depth of at least 10 μ m. According to a possible embodiment, the porous support contains fibrinogen in a layer which is at a distance of more than 10 μ m from the face in contact with the fibrin layer, particularly to a depth of 20 μ m.

The presence of free fibrinogen (having not reacted yet) has to be preferably avoided when the fibrin network has been already formed, in order to prevent new fibrin bonds from forming in the network upon reimmersion of the dried fibrin layer, such bonds reducing the size of the alveoli or of some alveoli of the network.

According to one embodiment of the invention, some of the fibrin attached to the network extends across the thickness of the treated porous part of the support, from said treated surface to a depth of at least 2 μ m, through the pores having an average diameter of 10 to 20 μ m and through the pores having an average diameter above 20 μ m. Although the support may be made of any hydrophobic or substantially hydrophobic material, it is particularly made of polyethylene, of polyethylene terephthalate or of polytetrafluoroethylene, said materials being advantageously stretched, particularly in both axial directions.

The term hydrophobic or substantially hydrophobic material is used herein to identify materials having a bias of 30 to 50°, which bias is measured with the method ASTM D 2578-84.

Advantageously, the porous part of the support has a substantially homogeneous and uniform porosity over the treated surface, i.e. the pore distribution or number by surface unit is substantially uniform for the porous part. For example, given one

porous part, the volume of the pores having a diameter of more than $10\mu\text{m}$ in an area of 1 cm^2 of the porous part varies from 0.8 to 1.2, preferably from 0.9 to 1.1 times the average volume of pores having a diameter of more than $10\mu\text{m}$, for each cm^2 of the porous part.

According to one embodiment, at least the face of the fibrin layer opposite to the one contacting the porous support is stabilized. Particularly, said fibrin-based layer is at least partially cross-linked, to form a network of adjacent alveoli, having apertures therebetween. The layer is advantageously sufficiently cross-linked not to be water-soluble. According to a detail of one advantageous embodiment, said layer is provided with cells and/or proteins, particularly with proteins mediating cell-fibrin bonds, with fibronectin, etc.

Although the thickness of the fibrin layer, when it is hydrated and re-hydrated may be of more than $100\mu\text{m}$, or even of more than $150\mu\text{m}$, according to a characteristic of one preferred embodiment, the cross-linked fibrin-based layer (in the hydrated or post-hydration state) which covers the porous part of the support is 0.5 to $100\mu\text{m}$ thick, advantageously 2.5 to $50\mu\text{m}$ thick, preferably 5 to $20\mu\text{m}$ thick, with alveoli being formed between the cross-linked fibrin-based molecules or bonds, said alveoli having a volume of 5 to $25\mu\text{m}^3$, the average thickness or height of said chamber being of 1 to $5\mu\text{m}$, particularly of 1 to $3\mu\text{m}$.

According to a detail of one particular embodiment, the pores of the support part, covered by said fibrin layer have inner faces at least partially covered by a water-soluble or substantially water-soluble protein. For example the pores of the support part covered by said fibrin layer are partially covered by fibrinogen, albumin, fibronectin, vibronectin, or by a mixture thereof. Particularly, the support face opposite to the treated face is at least partially

covered by a water-soluble or substantially water-soluble protein. Such covering is advantageous to improve the adhesion of tissues in contact with the face opposite to the treated face of the support.

5 In accordance with an advantageous characteristic, at least the pores of the porous part of the support are at least partially covered by a water-soluble or miscible polar additive. Such additive is preferably non-denaturing for protein and biocompatible
10 structures. Such additives may include glycerol, sugars (sucrose, mannitol, sorbitol, etc.). Said additives are particularly soluble or at least miscible in water and are particularly selected amongst water-soluble or
15 miscible additives allowing to lower the freezing temperature as compared with the water freezing temperature at atmospheric pressure.

According to a preferred embodiment, the element is dry, for example having a moisture content of less than 0.5% by weight, or even of less than 0.1% by
20 weight.

According to an advantageous embodiment, the fibrin layer is cross-linked in presence of fibronectin. The cross-linked fibronectin content in the fibrin layer is advantageously of 0.5 to 15%, preferably of 1 to 10%, of the fibrin and fibronectin weight in the
25 cross-linked layer. This content corresponds to the weight of fibronectin bonds in the layer as compared to the weight of fibrin and fibronectin bonds of the layer.

According to a detail of an advantageous
30 embodiment, the fibrin layer contains calcium, particularly calcium and chlorine, more precisely calcium chloride. The calcium content of the fibrin layer, expressed in μg of calcium by volume unit of the fibrin layer (cm^3) is advantageously of 1 to 100 $\mu\text{g}/\text{cm}^3$, preferably of 5 to 90 $\mu\text{g}/\text{cm}^3$, particularly of 10 to
35 50 $\mu\text{g}/\text{cm}^3$. The chlorine content in the fibrin layer is advantageously of 1.5 to 200 $\mu\text{g}/\text{cm}^3$, preferably of 8 to

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170 $\mu\text{g}/\text{cm}^3$, particularly of 16 to 100 $\mu\text{g}/\text{cm}^3$. When calcium is in the form of calcium chloride, the calcium chloride content in the fibrin layer (expressed in μg of calcium chloride by volume unit (cm^3) of the fibrin layer) is advantageously of 2.5 to 300 $\mu\text{g}/\text{cm}^3$, preferably of 13 to 260 $\mu\text{g}/\text{cm}^3$, particularly of 26 to 150 $\mu\text{g}/\text{cm}^3$.

5 Advantageously, the fibrin layer substantially contains no further salts of alkali or alkaline-earth metals in addition to calcium chloride.

10 Preferably, the content of salts of alkali or alkaline-earth metals differing from the calcium chloride is at least 10 times, preferably 20 times, particularly 50 times smaller than the content of calcium chloride in the fibrin layer.

15 Although the support may be a porous support whatsoever, the element support is preferably a biocompatible and/or biodegradable support.

20 According to a particular detail of one embodiment of the element in accordance with the invention, the element has two or more superposed fibrin layers. Advantageously, the layers have alveoli with different average volumes. Particularly, the fibrin layer which covers the fibrin layer in contact with the porous support has alveoli with a smaller average volume as compared with the average volume of the alveoli of the fibrin layer in contact with the porous support. For example, the average volume of alveoli in the fibrin layer which covers the fibrin layer in contact with the support is of less than about 0.5 times the average volume of alveoli in the fibrin layer in contact with the support. According to one embodiment, the fibrin layer covering the fibrin layer in contact with the support partially penetrates said fibrin layer in contact with the support. The penetration of the fibrin layer with small alveoli in the fibrin layer with large alveoli is advantageously such that the fibrin layer with small alveoli penetrates at least 50% of the

thickness of the fibrin layer with large alveoli, but preferably less than the whole thickness.

The fibrin of the layer of the element of the invention, as well the fibrin present in the porous substrate is substantially not denatured, preferably not denatured.

The invention also relates to a process for preparing an element according to the invention.

This process provides that:

- 10 - at least one porous part of a first face of a porous support is brought into contact with an aqueous solution containing fibrin or fibrinogen, or with one or more fibrin-based or fibrinogen-containing compounds,
- 15 - the face of the porous part of the support opposite to said first face is homogeneously and uniformly submitted to a suction force to suck the solution, at least partly, through said porous part, thus ensuring the deposition of a layer based on fibrin or on fibrinogen-containing materials, homogeneously and uniformly with respect to said porous part, and the diffusion of at least the solution water through the porous part of the porous support as well as the penetration of fibrin or fibrinogen-containing materials through the porous support. Such suction provides a fibrin layer which is substantially free of fibrinogen, particularly if the fibrin layer has been washed with water or with an aqueous solution. Advantageously, the suction of the solution through the porous material is carried out at least during the cross-linking of fibrin,
- 20 and preferably at least during the reaction of the fibrinogen-containing material and the cross-linking of the fibrin. The fibrin present in the porous material is therefore advantageously cross-linked with the fibrin layer covering the said first face of the porous material.
- 25
- 30
- 35

The process according to the invention provides an element which complies therewith, as described hereinbefore.

Thanks to suction, the fibrin attached to the network is arranged to penetrate the porous support to a depth of at least 2 μ m, both in the pores having an average diameter of 10 to 20 μ m and in the pores having an average diameter of more than 20 μ m.

Advantageously, the face of the support opposite to said first face, is submitted to a pressure of less than 0.8 10⁵ Pa, and a pressure difference is created between the two faces of the porous part of at least 0.3 10⁵ Pa. Preferably, the support face opposite to said first face is submitted to a pressure of less than 0.5 10⁵, preferably less than or equal to 0.4 10⁵ Pa. According to a preferred embodiment, while providing an efficient passage of fibrin or fibrinogen across the thickness of the porous part of the support, the support face opposite to the first face is intermittently submitted to a first pressure, of less than 0.8 10⁵ Pa, preferably less than 0.4 10⁵ Pa, said first pressure being at least 5% higher than the second pressure. Instead of varying the positive or negative pressure on the face opposite to said first face, it would be possible to slightly vary the pressure exerted on said first face.

Advantageously, the face of the porous support opposite to said first face is submitted to a pressure of less than 0.8 10⁵ Pa, and exposed to a temperature of 0 to 100°C, preferably to a temperature of 15 to 60°C, particularly to a temperature of 25 to 40°C.

According to a variant of the process according to the invention, the face of the porous support opposite to said first face is submitted to a solution selected so as to create a reverse osmosis, causing the diffusion of at least the solution water in

contact with the first face through the porous part of the support. Such diffusion ensuring thereby a substantially uniform and regular passage of fibrin or fibrinogen at least partially through the thickness of 5 the porous support.

For implementing the process of the invention, a solution containing fibrinogen-containing materials and thrombin is advantageously first prepared and after preparation contacted with the porous 10 material. Thereafter, the solution is at least partly sucked through the porous material. Preferably, substantially immediately after mixing the fibrinogen containing material and thrombin, the solution is contacted with the porous material and preferably sucked 15 through the porous material. For example, a solution containing fibrinogen containing material and thrombin is prepared in continue by mixing a solution of fibrinogen containing material with a thrombin containing solution, and after its preparation 20 (substantially immediately after its preparation), the said solution is contacted with the porous material. This is advantageous for ensuring a good distribution of the thrombin in the solution in contact with the porous material.

For implementing the process according to the invention, a solution is advantageously used which contains 5 to 20 mg/ml of fibrinogen-containing materials, particularly a solution which contains 5 to 30 20 mg/ml of fibrinogen-containing materials and 0.01 to 10 units of thrombin per ml, preferably a solution which contains 5 to 20 mg/ml of fibrinogen-containing materials, factor XIII, and 0.01 to 2, preferably 0.05 to 1 units of thrombin per ml. According to an advantageous embodiment, the solution contains less than 35 0.5 units of thrombin per ml.

Advantages have also been noted with a solution containing 0.1 to 10 units of factor XIII per

ml. Advantageous results have also been obtained from a solution containing 1 to 40 millimoles of CaCl_2/ml , particularly 1 to 20 millimoles of CaCl_2/ml to reduce or slow down fibrinolysis. Hence, for example, for a fibrin 5 layer prepared with 20 millimoles of CaCl_2/ml , no fibrinolysis was visually detected one week after the fibrin layer had been prepared.

It will be noted that smaller quantities of thrombin used in the formation of the fibrin network 10 correspond to larger amounts of fibrinogen which can penetrate the porous support. In spite of this, the process according to the invention provides a fibrin layer substantially free of fibrinogen, particularly at the face of the support which is in contact with the fibrin layer.

According to a characteristic of a process according to the invention, during a first step, at 15 least one portion of a first face of a porous support is placed into contact with a solution containing fibrin and/or fibrinogen-containing materials, while the face of the porous support opposite to said first face is homogeneously and uniformly submitted to a suction force, thus ensuring a diffusion of at least the solution water across the thickness of the porous support and a penetration of fibrin or fibrinogen-containing materials in the porous support to a depth of 20 at least $2\mu\text{m}$, homogeneously and uniformly with respect to said porous part of the first face and, during a second step, the fibrin and/or fibrinogen layer is 25 stabilized.

In accordance with a possible embodiment, a contact is provided between said part of the first face and a moving aqueous solution.

Advantageously, the solution containing 30 fibrin or materials containing fibrinogen also contains a polar organic additive. The use of such polar organic additive has proved to allow the control of fiber

thickness in the fibrin network. Moreover, the presence of such organic additive has also provided advantages in the protection of the fibrin-based layer during the drying step, which may be possibly provided after a 5 washing step. The drying operation is advantageously effected at least partially by lyophilization, advantageously at a temperature of -30°C and -100°C, preferably at a temperature of -40°C to -70°C. For example, the drying operation is performed in several 10 steps, i.e. a first drying step for raising temperature (for example at a temperature of 30 to 70°C) or for creating a vacuum after removal of the fibrin or fibrinogen-containing material solution in contact with the porous part of the support, and a second drying step 15 for lyophilization.

Drying operations are advantageously performed after one or more washing steps, by means of water, an aqueous solution, e.g. an aqueous solution containing a polar organic additive (e.g. in the order of 1 to 20% by weight, particularly in the order of 5 to 20 10% by weight), such as glycerin. A particular washing operation consists in bringing the fibrin layer integral with the porous support in contact with an aqueous solution, particularly a solution containing glycerol 25 (e.g. 1 to 20% by weight, particularly 5 to 10% by weight) and thereafter in submitting the other face of the support to a suction force, to suck the solution through the porous support. Such washing operation provides a fibrinogen-free porous support. This 30 operation may be performed on supports provided with a fibrin layer which are not compliant with the invention, thereby allowing to turn a product obtained by a simple contact of the porous support with the fibrinogen-containing solution into an element according to the 35 invention.

The solution of fibrin or of fibrinogen-containing materials used in the process for forming the

fibrin layer according to the invention preferably contains 0 to 20%, particularly 3 to 15%, and more particularly 5 to 10% of said polar organic additive. This additive may advantageously be glycerol, a sugar (mannitol, sorbitol, sucrose, glucose, etc.). When using a solution which contains fibrinogen-containing materials in the order of 5 to 20 mg/ml, thrombin in the order of 0.01 to 10 units/ml and 5 to 10% of glycerol in the process according to the invention, a network of fibrin fibers was obtained, wherein the size of the alveoli is similar to that in the network obtained with a solution which contains fibrinogen-containing materials in the order of 5 to 20 mg/ml, thrombin in the order of 0.01 to 10 units/ml (without glycerol) in the process according to the invention. Nevertheless, the fiber size in the network obtained by using glycerol was smaller, whereby a better use of fibrin or fibrinogen-containing materials in the solution resulted when using glycerol.

The pH of the solution of fibrin or of fibrinogen-containing materials is advantageously of 5 to 8.5, preferably of 5.5 to 8, particularly of 6 to 7.5. The pH of the solution may be controlled by means of a buffer solution (e.g. a tris buffer), by adding a strong (HCl) or weak acid, of mineral or organic origin (citric acid, etc.).

The solution also advantageously contains at least a water-soluble protein, particularly albumin.

According to one particular embodiment, at least for a part of the deposit of the fibrin- or fibrinogen compounds-based layer, the concentration of fibrin or fibrinogen compounds in the solution in contact with the first face is controlled in order to ensure a substantially constant water diffusion through the support.

In the process according to the invention, a biocompatible or biodegradable porous support is used.

According to a particular embodiment, wherefrom advantages are obtained to ensure from the start a substantially uniform water diffusion through the thickness of the porous support, the porous part is treated with an aqueous solution which advantageously contains a wetting agent and/or a water-soluble protein and/or a polar organic additive, before bringing the fibrin- or fibrinogen-containing solution in contact with said porous part.

According to the invention, the porous support may be also treated, successively, with a solution which contains fibrin or fibrinogen-containing materials to deposit several fibrin layers. According to the invention, the porous support may be treated with a solution which contains fibrin or fibrinogen-containing materials but does not contain thrombin, and then the pretreated support may be treated with a solution containing thrombin.

The invention also relates to a filter including a filtering membrane consisting of an element according to the invention, to a bioreactor including a membrane consisting of an element according to the invention, an implant consisting of an element according to the invention, and an artificial skin produced from an element according to the invention.

Since glycerol has been found to be useful for controlling the size of alveoli, for a better use of fibrin (thinner fibers) and for ensuring a better viability of the cells attached to the fibrin network, another object of the invention is a compound based on fibrin or on fibrinogen-containing materials, said compound having the form of a dry foam or of particles of dry foam, containing 0.05 to 10% by weight of a water-soluble or miscible polar organic additive, said foam having a porosity consisting of at least 50% by volume of chambers or volume cavities of 5 to 25 μm^2 . Advantageously, at least 90% by weight of fibrin is in

cross-linked form. Possibly, the compound also contains one or more proteins and/or one or more active substances. Amongst polar additives, glycerin is preferred, but other additives may be also used, such as 5 sugars, sucrose, glucose, mannitol, etc. The water content is advantageously lower than 0.5% by weight. In fact the foam or cross-linked fibrin network is at least partially covered by a water-soluble or miscible polar organic additive.

10 The preparation of this compound may be effected in a process wherein, possibly after a pre-drying step, an aqueous solution of fibrin and/or fibrinogen, also containing a water-soluble or miscible polar organic additive, is dried by lyophilization, the 15 organic solvent content of said solution being of 0.05 to 10% by weight, so as to obtain a compound containing less than 0.5% thereof by weight. Advantageously, the drying operation by lyophilization is effected at a temperature of -40 to -100°C, preferably of -50°C to 20 -75°C. Particularly, lyophilization is performed in three steps, each step involving a temperature decrease of the compound or solution to a temperature of -40 to -100°C, followed by a pressure decrease to less than 0.4 bar ($0.4 \cdot 10^5$ Pa). For example, in a first step, pressure 25 is lowered to a pressure of 0.2 to $0.4 \cdot 10^5$ Pa, and in the last step, pressure is decreased to less than $0.2 \cdot 10^5$ Pa.

30 The invention also relates to a process allowing to extract the unbound fibrinogen from the fibrin layer, and particularly the fibrinogen which is found in the porous support, in such a way as to obtain a fibrinogen-free fibrin layer, and particularly a porous support and a fibrin layer both free of fibrinogen. This process provides that:

35 - at least one part of the fibrin layer attached on a first face of the porous support is brought to contact with an aqueous solution advantageously containing a polar organic additive, and

- the face of the porous support opposite to said first face is homogeneously and uniformly submitted to a suction force to suck the solution, at least partly, through said porous part, thus ensuring the removal of fibrinogen in the proximity of said first face of the support, homogeneously and uniformly with respect to said porous part. Thanks to this suction, at least the solution water is diffused through the thickness of the porous part of the porous support. If this process is applied for a sufficient time, the amount of water diffused through the thickness of the porous part can be sufficient to remove or extract the fibrinogen in the pores of the porous support. Therefore, this suction provides a substantially fibrinogen-free fibrin layer, or even a porous support and a fibrin layer free of fibrinogen.

This washing process, when using an aqueous solution which contains one or more additives, e.g. one or more soluble proteins, one or more drugs, etc, allows the introduction in the porous support of a certain amount of said additive/s, or the covering of the face of the support which is not in contact with the fibrin layer with said additive/s.

Thanks to the solution suction, water is allowed to penetrate the porous support, so that, for example, at a depth of at least 2 μm from the first face (face bearing the fibrin layer), advantageously of at least 10 μm , preferably of at least 20 μm , at least the pores having an average diameter of 10 to 20 μm are free of fibrinogen.

Advantageously, the face of the porous support opposite to said first face is submitted to a pressure of less than $0.8 \cdot 10^5 \text{ Pa}$, and a pressure difference of at least $0.3 \cdot 10^5 \text{ Pa}$ is created between the two faces of the porous part. Preferably, the face of the porous support opposite to said first face is submitted to a pressure of less than $0.5 \cdot 10^5 \text{ Pa}$, more

preferably less than or equal to $0.4 \cdot 10^5$ Pa. According to a preferred embodiment, while providing an efficient passage of water across the thickness of the porous part of the support, the face of the porous support opposite to the first face is intermittently submitted to a first pressure of less than $0.8 \cdot 10^5$ Pa, preferably less than about $0.4 \cdot 10^5$ Pa, and to a second pressure of less than $0.8 \cdot 10^5$ Pa, preferably less than $0.4 \cdot 10^5$ Pa, the first pressure being at least 5% higher than the second pressure. Instead of varying the positive or negative pressure on the face opposite to said first face, it would be possible to slightly vary the pressure exerted on said first face.

Advantageously, the face of the porous support opposite to said first face is submitted to a pressure of less than $0.8 \cdot 10^5$ Pa, and to a temperature of 0 to 100°C, preferably to a temperature of 15 to 60°C, particularly to a temperature of 25 to 40°C.

According to a variant of the process according to the invention, the face of the porous support opposite to said first face is submitted to a solution selected so as to create a reverse osmosis, causing the diffusion of at least the solution water in contact with the first face through the porous part of the support. Such diffusion ensuring thereby a substantially uniform and regular passage of water at least partially through the thickness of the porous support.

A further object of the invention is a process for preparing porous supports covered by a layer made of a bioabsorbable material or of an absorbable polymer, particularly of a polylactic polymer and/or of polyglycol polymers and/or of biopolymers, as well as structural proteins and polysaccharides, said structural proteins being selected in the group including collagen, elastin, fibronectin, laminin and fibrin, and other proteins forming human or animal tissues, as well as

recombinant proteins. This process provides that an aqueous solution or suspension is prepared, which contains one or more polymers and/or biopolymers and/or materials to form said polymers and/or biopolymers on site. This solution or suspension is brought to contact with a first face of a porous support, while sucking at least a part of the water of said solution or suspension from at least one different face of the porous support (advantageously the face opposite to the first face). This suction force causes water and advantageously absorbable biopolymers or polymers to be diffused in the porous support. In order to ensure such diffusion, the face of the porous support opposite to said first face (face in contact with the solution or suspension) is submitted to a pressure of less than $0.8 \cdot 10^5$ Pa, while a pressure difference is created between the two faces of the porous part of at least $0.3 \cdot 10^5$ Pa. Preferably, the support face opposite to said first face is submitted to a pressure of less than $0.5 \cdot 10^5$ Pa, preferably less than or equal to $0.4 \cdot 10^5$ Pa. According to a preferred embodiment, providing an efficient passage of water through the thickness of the porous part of the support, the face of the support opposite to the first face is intermittently submitted to a first pressure, of less than $0.8 \cdot 10^5$ Pa, preferably less than $0.4 \cdot 10^5$ Pa, and to a second pressure, of less than $0.8 \cdot 10^5$ Pa, preferably less than $0.4 \cdot 10^5$ Pa, the first pressure being at least 5% higher than the second pressure. Instead of varying the positive or negative pressure on the face opposite to said first face, it would be possible to slightly vary the pressure exerted on said first face. Said face opposite to the face in contact with the polymer solution or suspension might also be submitted to the influence of a solution selected so as to create a reverse osmosis, causing the diffusion of at least the solution water in contact with the first face through the porous part of the support. Such diffusion ensuring

thereby a substantially uniform and regular passage of water at least partially through the thickness of the porous support. The solution diffusing through the porous support advantageously is at a temperature of 20 to 70°C, particularly of 30 to 50°C. Once the layer of absorbable polymers or biopolymers is formed, this layer is advantageously dried by lyophilization. Lyophilization is advantageously effected as described with respect the fibrin layer. If drying operations are performed by lyophilization, the solution used to form the layer advantageously contains a polar additive, particularly glycerol, for example in the order of 1 to 15%, particularly of 5 to 10%.

Further characteristics and details will be apparent from the following detailed description of certain embodiments, wherein reference is made to the annexed drawings. In these drawings,

- figure 1 is a diagrammatic view of an element according to the invention;

- figure 2 is a diagrammatic view of an installation for preparing an element according to the invention;

- figures 3, 4 and 5 are cross-sectional views of a slice of fibrin networks, as taken with an electron microscope (Philips XL20 Scanning Electron Microscope), with a magnification of 5,000 times, before lyophilization, whereas figures 6, 7 and 8 are cross-sectional views of a slice of fibrin networks, as taken with an electron microscope, with a magnification of 5,000 times, after lyophilization;

- figures 9, 10 and 11 are cross-sectional views of the networks obtained by means of a solution containing 1 IU/ml, 10 IU/ml and 20 IU/ml of thrombin respectively, as seen in cross section;

- figures 12 to 14 are cross-sectional views of networks obtained by means of a solution containing no CaCl_2 (figure 12), 2.7 mM CaCl_2/ml (figure 13) and

27 mM CaCl_2/ml (figure 14), as taken with an electron microscope (Philips XL20), with a magnification of 5,000 times;

5 - figures 15, 16, 17 and 18 are top views of the fibrin networks with cells after two hours of culture, as taken with an electron microscope (Philips XL20 Scanning Electron Microscope), with a magnification of 500 times;

10 - figures 19 to 21 are cross sectional views of a porous support 2, bearing a fibrin layer 4, with cells "C", as taken with an electron microscope, with a magnification of 100 times, 100 times and 1000 times respectively;

15 - figure 22 is an electrophoresis diagram of markers having a low molecular weight (1, 6), of control fibrinogen (5), of control fibrin (4), of the polymer layer from the exudate (the part passing through the porous membrane) after incubation, and of the floating part of the exudate after incubation.

20 Figure 1 diagrammatically shows a sectional, larger-scale view of one part of an element according to the invention.

25 The element 1 comprises (a) a hydrophobic or substantially hydrophobic support 2, for example PTFE (expanded and stretched in both axial directions), which has a porous part with a thickness E of 0.1 to 5 mm, e.g. of 300 to 500 μm , and whose pores P , extending across its thickness have an average diameter "d" (porous volume/surface of pores) of 5 to 100 μm , e.g. of about 30 to 40 μm , one face 3 of said porous part of said support 2 being treated with a fibrin- and/or fibrinogen-based compound, and (b) a fibrin-based layer 4 covering said treated surface 3 of the support 2.

30 Said fibrin-based layer 4 is substantially uniform and homogeneous on said treated face 3. After being washed, the fibrin layer 4 contains no fibrinogen. For example, the content of fibrinogen in the layer 4

(fibrinogen unbound from the fibrin layer) is below 0.5% by weight; preferably below 0.1% by weight of the fibrin layer.

Some fibrinogen F may extend across the thickness E of the treated porous part of the support, from said treated face to a depth "e" of at least 10 μm both in the pores having an average diameter of 10 to 20 μm (pores whose volume, expressed in μm^3 , divided by the surface of the pore walls, expressed in μm^2 , gives 10 to 20 μm) and in the pores having an average diameter of more than 20 μm . Particularly, in all the pores of more than 25 μm of the treated face of the porous part, some fibrinogen extends across the thickness E of the support to a depth "e" of at least 30 μm . Nevertheless, at the face 3, the support is substantially free of fibrinogen unbound from the network. The lack of fibrinogen unbound from the fibrin network is due to the passage of water through the porous support. In one particular embodiment, the porous support is free of fibrinogen to a depth of at least 10 μm , from the face bearing the fibrin layer. According to a particularly advantageous embodiment, the support is free of fibrinogen throughout its thickness.

The fibrin layer 4, as shown in figure 1, is stabilized by cross-linkage due to the presence of factor XIII. Hence, said layer 4 forms a network of adjacent alveoli 40.

The thickness "h" of the fibrin layer as determined from the face 3 (in its dehydrated form) is, for example, of about 10 μm , while the average volume of a chamber or cell is of the order of 10 μm^3 . The alveoli are open and have apertures therebetween. The term alveoli defines fibrin-free areas having a volume of more than 5 μm^3 , surrounded by fibrin bonds. The distribution of alveoli over the layer 4 is substantially regular, that is the volume of the alveoli over an area of 1 cm^2 of the face 3 covered by the layer

4 is of 0.8 to 1.2 times (preferably of 0.9 to 1.1 times) the average volume of chambers by unit of surface (cm²) of that area. The average height of the chambers, as measured perpendicularly to the face 3 is, for example, of 2 to 3 µm.

The element as shown in figure 1 is advantageously in a dry state. The moisture content is, for example, of less than 0.01% by weight, which ensures an excellent preservation and stability of the element. When the element is rehydrated, the fibrin layer inflates, for example, by a factor of more than 1.5, particularly by a factor of 1.6 to 2.5 (the thickness of the fibrin layer after rehydration corresponds to 1.6 to 2.5 times the thickness of the dry fibrin layer).

According to a particular embodiment, the pores P have inner faces at least partially covered by a water-soluble or substantially water-soluble protein and/or the face 6 of the support, opposite to the treated face is at least partially covered by a water-soluble or substantially water-soluble protein. Such covering is advantageous to assist, for example, cell fixation, the adhesion of the tissues surrounding the face opposite to the face treated with fibrin or with fibrinogen-containing materials.

According to an advantageous characteristic of one embodiment, the pores P (inner walls) of the porous part of the support are at least partially covered by a water-soluble or miscible polar organic additive or by traces of such additive. This polar organic additive is advantageously also present at least in part on the fibrin layers of the layer 4 and on the faces 3 and 6 of the support. This additive may be, for example, glycerol, a sugar, etc. or a mixture of these additives. Said additives are particularly soluble or at least miscible in water and are particularly selected amongst water soluble additives allowing the freezing temperature to be lowered with respect to water

freezing temperature at atmospheric pressure. The amount of soluble or miscible additive in the fibrin, fibrinogen and/or thrombin solution or in the wet cross-linked fibrin layer (not dried, the water content in the pores is in the order of 50%) is preferably sufficient to lower the freezing temperature at atmospheric pressure of less than -5°C, preferably of less than -10°C.

Although the support of the illustrated embodiment is a biocompatible porous support of PTFE, another biocompatible support can be used, particularly a biodegradable support, or a biocompatible and biodegradable support.

A few examples of processes for preparing an element according to the invention will be described hereafter.

For the preparation of one or more elements according to the invention, a vacuum chamber 10, connected to a vacuum pump 11 has been used to create a vacuum or a negative pressure in the chamber with respect to atmospheric pressure. This chamber is shown diagrammatically in figure 2.

The chamber has an intake for letting the solution/s into the inner space or hollow of a tube having an inside diameter of 1 to 100 mm, more particularly of 2 to 10 mm. The tube 13 has porous cylindrical parts 13A (average diameter of pores of 20 to 30 μm) separated by a non-porous ring 13B. The tube thickness was of about 200 to 300 μm for the porous parts. The intake 12 includes the means of fastening an end 13C of the tube thereto in a fluid-tight manner. The intake 12 is connected by means of a duct 15 to a tank 14 which contains an aqueous solution of a fibrinogen-containing material (with a concentration of 10 to 40 mg/ml), including 0.2 to 20 units of factor XIII per ml (IU/ml) and 100 to 1000 $\mu\text{g}/\text{ml}$ of fibronectin, by means of a duct 16 to a tank 17 which contains an aqueous

solution of thrombin (with a concentration of 0.05 to 2 IU/ml) and by means of a duct 18 to a tank 19 which contains water and possibly one or more additives. The ducts 15, 16 and 18 are fitted with valves ν to allow or prevent the passage of a fluid. Said ducts lead one or more fluids towards the intake, depending on atmospheric pressure. A control system 20 controls the vacuum pump operation depending on the desired vacuum and on the vacuum measured inside the chamber.

The tube end opposite to the one fastened to the intake is closed by a plug 21, advantageously extended by a duct 22 with a valve 23, to allow the evacuation of fluids or solutions contained in the tube.

The chamber is also provided with a means 24 to adjust the chamber temperature in the range of +60°C to -100°C.

Example 1

In this example, a solution A, containing 20 mg/ml of a fibrinogen-containing material, 1000 μ g of fibronectin per ml and 2IU/ml of factor XIII, and a solution B, containing 0.2IU of thrombin per ml, and 40 mM (millimoles) of calcium chloride per ml, were used.

The solution A and the solution B were fed into the intake at the same flow rate to obtain a 1:1 mixture of both solutions A and B. The mixture obtained thereby contained 10 mg/ml of fibrinogen, 500 μ g/ml of fibronectin, 1 IU/ml of factor XIII, 0.1 IU/ml of thrombin and 20 mM/ml of CaCl_2 .

The hollow or inner space of the tube was filled with the mixture of solutions A and B, and the chamber pressure was lowered to $0.4 \cdot 10^5$ Pa (that is a negative pressure of about 600 millibar with respect to atmospheric pressure). This vacuum creation causes water to be sucked in through the thickness of the porous parts of the tube. Since the vacuum is created on the outer surface of the tube, the latter is slightly

stretched or tightened, which assists the diffusion of liquid through the pores of the tube.

While creating and maintaining vacuum, the outer wall of the tube was found to be wet.

After maintaining the vacuum for about 1 to 30 minutes, the chamber pressure was progressively reset to atmospheric pressure. Once the tube was emptied and washed with water, the inner face of the tube was found to be covered by a cross-linked fibrin layer about 5 μm thick, with chambers or open cells of 15-20 μm^3 on the porous parts of the tube (cell height of 2 to 3 μm , area of 5 to 7 μm^2 , as measured parallel to the face of the support bearing the layer). No fibrinogen unbound from the fibrin layer was found in the fibrin layer, nor on the support interface with the fibrin layer. Fibrinogen was found in the pores of the support to a depth (from the inner surface of the tube) of at least about 20 μm for all pores having an average diameter of more than 10 μm .

The passage of fibrinogen through the porous support is confirmed by the electrophoresis diagram of figure 22. In fact, some liquid from the face opposite to the one in contact with the fibrinogen solution was collected. After incubating this liquid, electrophoresis peaks were determined both for the polymer layer formed (2) and for the supernatant (3). The result was that, after incubation, the electrophoresis (2) showed fibrin-typical peaks, which proves that fibrinogen had passed through the porous support.

This tube was subsequently dried by a gas heated to 50°C.

Example 2

Example 1 was repeated, except that the washing step was effected by letting demineralized water flow inside the tube to evacuate the fibrinogen solution, while maintaining a pressure of about

0.4 10⁵ Pa in the chamber to ensure a diffusion of washing water through the porous support. This diffusion allows fibrinogen to be removed from the porous support.

Example 3

Example 1 was repeated, except that the tube was dried by lowering the tube temperature to -60°C to turn water into ice and by lyophilizing it at this temperature.

Example 4

Example 3 was repeated, except that glycerol was added in the order of 5% by weight of the mixture consisting of 50% of the solution A and 50% of the solution B. It was noted that the presence of glycerol both in the porous support and in the fibrin layer provided a certain flexibility of the element. Further, the lyophilization step was easier.

The presence of glycerol upon formation of the cross-linked fibrin proved to be advantageous for providing a regular and homogeneous structure of the fibrin layer. Moreover, the presence of glycerol assisted the passage of fibrin and fibrinogen in the pores of the porous part of the tube.

Example 5

Example 4 was repeated, except that glycerol was added in the order of 10% by weight of the mixture consisting of 50% of solution A and 50% of solution B. It was noted that the presence of glycerol both in the porous support and in the fibrin layer provided a certain flexibility of the element. Further, the lyophilization step was easier.

Some parts of the fibrin networks from examples 2 and 4, before and after lyophilization were left for one night in dishes containing a solution of 2 to 2.5% of glutaraldehyde in dishes. Thereafter, a slice of the network fixed by glutaraldehyde was cut transversely by means of a heated scalpel, which slice

was dehydrated by 40%, 50%, 70%, 80%, 90% and 100% ethanol solutions.

Figures 3, 4 and 5 are cross-sectional views of slices of fibrin networks from examples 2, 3 and 4 respectively, as taken with an electron microscope (Philips XL20 Scanning Electron Microscope), with a magnification of 5,000 times, before lyophilization, whereas figures 6, 7 and 8 are cross-sectional views of slices of fibrin networks from examples 2, 3 and 4 respectively, as taken with an electron microscope, with a magnification of 5,000 times, after lyophilization. By comparing these figures, the result is that the alveoli of the fibrin network from examples 2, 3 and 4 before lyophilization are similar, that the alveoli of the fibrin network from examples 2, 3 and 4 after lyophilization are similar, and that the use of glycerol allows the size of the network fibers to be reduced. Hence, glycerol, besides being useful to protect fibers during the lyophilization step, is an agent allowing control of the size or the diameter of the fibers of the fibrin network.

Example 6

Example 4 was repeated, except that glycerol was replaced first by glucose and then by mannitol.

Example 7

Example 1 was repeated, except for the use of a solution containing fibrinogen in the order of 10 mg/ml and thrombin in the order of 1IU/ml, 10IU/ml and 20 IU/ml respectively.

The networks obtained thereby were treated with a solution containing 2 to 2.5% of glutaraldehyde and with ethanol-containing solutions as described in example 4. Some slices of the networks so obtained were examined with an electron microscope (scanning electron microscope, Philips XL20). Figures 9, 10, and 11 are cross sectional views of the networks obtained with a solution containing 1 IU/ml, 10 IU/ml and 20 IU/ml of

thrombin respectively, with a magnification of 3,500 times.

These figures 9 to 11 show that a higher concentration of thrombin in the solution produces a greater number of fibers, but a smaller size thereof.

s Example 8

Example 1 was repeated, except that thrombin and fibrinogen solutions were prepared, which had a CaCl_2 concentration of 0mM/ml, 2.7mM/ml and 27 mM/ml. After 10 treating and washing the networks as described in example 4, the cross section of the networks obtained with a solution containing 0mM/ml (figure 12), 2.7mM/ml (figure 13) and 27 mM/ml (figure 14) was examined with an electron microscope (Philips XL20), with a 15 magnification of 5,000 times. These figures show that a higher calcium content corresponds to a greater number of fibers, a larger size thereof, and a smaller volume of the alveoli.

Example 9

Example 4 was repeated, except for the use of 20 a solution containing 5% of glycerol, 20 mg/ml of fibrinogen, 500 μg of fibronectin per ml, 10 IU/ml of factor XIII, 1 IU of thrombin per ml and 40 mM (millimoles) of calcium per ml.

Example 10

Example 4 was repeated, except that the 25 chamber vacuum was controlled to cause its intermittent variation from 600 mbar with respect to atmospheric pressure (a pressure of about $0.4 \cdot 10^5 \text{ Pa}$) to 630 mbar with respect to atmospheric pressure (a pressure of about $0.38 \cdot 10^5 \text{ Pa}$). This vacuum variation was found to be 30 advantageous for fibrin and fibrinogen diffusion in the pores of the support. After washing with water, bringing the fibrin layer in contact with a water flow and 35 creating a vacuum in the chamber varying from 600 mbar with respect to atmospheric pressure (a pressure of about $0.4 \cdot 10^5 \text{ Pa}$) to 630 mbar with respect to

atmospheric pressure (a pressure of about $0.38 \cdot 10^5$ Pa), the support and the fibrin layer contained no more free fibrinogen.

The tube may be easily sterilized, if needed, before or after lyophilization, at a temperature of 121°C for 60 minutes, for example in an autoclave. Any other sterilization method may be used, provided that it does not destroy the alveoli structure of the cross-linked fibrin layer, nor the support structure.

10 Example 11

Example 3 was repeated, except that the fibrinogen concentration was controlled in the tube, during the diffusion step, so as to ensure a substantially constant fibrinogen concentration in the tube. In order to do this, valve 23 was intermittently opened to evacuate a certain amount of solution out of the tube and a solution containing little or no fibrinogen was fed into the tube. This ensures that the fibrin layer is substantially regular and homogeneous in thickness.

15 Example 12

Example 11 was repeated, except that the fibrinogen concentration was controlled substantially continuously, to decrease this concentration as fibrin is deposited on the inner wall of the tube.

20 Example 13

Example 3 was repeated, except that, before treating the tubes with the fibrinogen solution, demineralized water, an aqueous solution containing 1 mg/ml of albumin, an aqueous solution containing 10 mg/ml of albumin, an aqueous solution containing 30 mg/ml of albumin, were respectively fed into the tubes, so as to fill or saturate the pores with said solution, before treating the tubes with the fibrinogen solution.

25 Example 14

Example 3 was repeated, except that the proteins contained in the solution were 30 mg/ml of

albumin and 10 mg/ml of fibronectin. Other proteins, such as vibronectin, etc. could be used, individually or in mixture, instead of albumin and/or fibronectin.

5 As set out in WO96/07444, the fibrin layer can be treated either to denature it or to provide it with particular properties.

10 The fibrin layer may be treated with water, with one or more salts (possibly in solution), with additives used to improve the biocompatibility of the support provided with the fibrin layer. The additives may be selected, for example, amongst proteins, 15 anticoagulants, anti-inflammatory compounds, compounds reducing graft rejection, living cells, cell growth inhibitors, agents stimulating endothelial cells, antibiotics, antineoplastics, genetic materials, proteins promoting or stimulating the growth and/or attachment of endothelial cells on the cross-linked fibrin layer, growth factors, growth factors for heparin bond, substances against cholesterol (ZOCOR®), etc. Some 20 particular examples of additives are given in US 5,660,873, whose content is included in this application by way of reference.

25 The fibrin layer may be partially hydrolyzed, if needed, for example by means of a plasmin.

Example 15

30 Example 1 was repeated, except that, during a first step, solution A was fed into the tube to obtain, by creating vacuum in the chamber, a non-cross-linked fibrin or fibrinogen layer, and that, during a second step, solution B (thrombin) was fed into the tube to form fibrin monomers and to obtain a cross-linked structure.

Example 16

35 Example 4 was repeated, except that lyophilization was effected in several steps, i.e. by lowering temperature to -58°C, by maintaining this

temperature of -58°C, by creating a vacuum (the lyophilization device had been adjusted with a pressure set-point of 7 Pa, so that the vacuum pump could operate continuously) for 1 to 5 hours, by raising temperature 5 from -58°C to -20°C to -30°C, while maintaining the vacuum, by maintaining the temperature of -20°C to -30°C, while maintaining the vacuum, for at least 10 hours (10 to 100 hours), by increasing the temperature to more than 20°C, while maintaining the vacuum.

10 Example 17

Example 1 was repeated, except that the porous tube was successively treated with solution A and with solution B.

15 The treatment steps of this example are:

- a) feeding solution A (fibrinogen) into the tube;
- 15 b) creating a vacuum in the space outside the tube to suck solution A through the walls of the tube;
- c) removing solution A still present in the 20 tube;
- d) incubating the fibrinogen layer deposited for 15 minutes at ambient temperature (steps a), b), c) and d) may be repeated once or several times, for example twice or three times before step e));
- 25 e) feeding solution B (thrombin) into the tube;
- f) creating a vacuum in the space outside the tube to suck solution B through the walls of the tube;
- 30 g) removing solution B still present in the tube;
- h) incubating the layer at 37°C for 30 minutes;
- i) feeding solution A (fibrinogen) into the tube;
- 35 j) creating a vacuum in the space outside the tube to suck solution A through the walls of the tube;

k) removing solution A still present in the tube;

5 s 1) incubating the layer for 15 minutes at ambient temperature (steps i, j, k, and l may be repeated once or several times);

m) incubating the layer for 90 minutes at 37°C.

Example 18

10 Example 16 was repeated, except that the intermediate incubation steps d, h and l were skipped.

Example 19

15 Example 1 was repeated, except that the porous tube was successively treated with solution A and with solution B.

15 The treatment steps of this example are:

a) feeding solution A (fibrinogen) into the tube;

b) creating a vacuum in the space outside the tube to suck solution A through the walls of the tube;

20 c) removing solution A still present in the tube;

d) incubating the fibrinogen layer deposited for 15 minutes at ambient temperature;

25 e) feeding solution B (thrombin) into the tube;

f) creating a vacuum in the space outside the tube to suck solution B through the walls of the tube;

g) removing solution B still present in the tube;

30 h) incubating the layer at 37°C for 30 minutes;

i) washing the tube with water (preferably in successive washing operations);

35 j) steps a to i are repeated once or several times;

k) incubating the layer for 90 minutes at 37°C.

Example 20

Example 4 was repeated, except that the pH of the solution mixture was changed, upon its introduction, to 6, 6.5, 7, and 7.5 respectively, or except that the pH of the solution in the tube was controlled during the process to maintain it, for example, at 6.5 or 7 or 7.5.

Example 21

Example 1 was repeated, except that, instead of placing the porous tube in a vacuum chamber, the tube was placed in a container with a concentrated aqueous solution of salt (NaCl) in order to create, by reverse osmosis, a water and fibrin-fibrinogen diffusion through the wall of the tube towards said concentrated solution.

Example 22

The fibrinogen and thrombin compound of example 1 was injected by means of a syringe in a tube, to create a fibrin layer on the inner wall of the tube. This process causes fibrinogen to be present on the inner wall of the tube and in the fibrin layer in the proximity of said inner wall.

After removal of the fibrinogen solution and immersion of the tube in water (prewashing) the tube was placed in the vacuum chamber used in example 1. Then, demineralized water was fed into the tube, whereupon a vacuum was created in the chamber (pressure of $0.3 \cdot 10^5$ Pa), so that water is sucked through the wall of the tube from the inner wall to the outer wall. This diffusion of water through the tube wall allows the unbound fibrinogen to be removed from the fibrin layer and outside the support, so that at least the part of the tube situated in the proximity of the inner wall of the tube is free of fibrinogen.

Example 23

Example 22 was repeated, except that an aqueous solution containing 5% of glycerol was used for the washing operation by diffusion of water through the tube wall.

Example 24

Example 22 was repeated, except that an aqueous solution containing 5% of glycerol and 1% of albumin was used for the washing operation by diffusion of water through the tube wall.

In the above examples, fibrin layers were prepared by using fibrinogen and thrombin from human blood. These could be replaced by products available on the market, such as biological glues by CRYOLIFE, e.g. the product FibRx, or by VITEX (the product VIGuard), or even recombinant fibrinogen.

The elements or membranes according to the invention, for example the membranes of examples 1 to 13 may be used in several applications, namely as membranes for bioreactors, for example as described in the application EP 96910867, as membranes for filters, as implants such as artificial internal organs, as artificial veins, as artificial arteries, as antithrombotic materials, as cardiac valves, as artificial skins; the membrane may also be applied to the production of test kits or equipment, etc...

A number of tests was performed to determine the morphology of the cells attached to a lyophilized fibrin network prepared with no added glycerol (example 3), to a fibrin network prepared with a solution containing about 5% of glycerol (example 4) before and after lyophilization, and to a fibrin network prepared with a solution containing about 10% of glycerol (example 6) with lyophilization.

In these tests, a culture medium was prepared, from Dulbecco Modified Eagle Medium (DMEM). The following components, in the weight % as specified hereafter, were added to this DMEM medium:

- 20% of HAM'S F 12 (culture medium)
- 10% of FCS (Foetal Calf serum)

- 1% of non essential amino acids (i.e. L-alanine, L-asparagine, L-aspartic acid, L-glutamic acid, Glycine, L-proline, L-serine)
- 1% of sodium pyruvate
- 1% of *Penicillium streptomycin*, and
- 1% of L-glutamine.

5 This medium will be hereafter termed "prepared DMEM medium".

10 The cells used in these tests were isolated as follows:

15 Just after the slaughter of cows, the bovine aorta was recovered. After separating the adipose tissues of the aortas, the collateral arteries were ligatured. The inner surface of the aortas was treated for 15 minutes at 37°C with a solution containing 250 IU/ml of collagenase. The cells released in this treatment were recovered and placed in a DMEM culture medium containing valine D, 10% of FCS, 100 IU/ml of penicillin, 100 µg/ml of streptomycin and 2.5 µg/ml of amphotericin B. The culture medium was renewed after 24 hours.

20 25 After two days, the culture medium was placed in a 70% DMEM culture medium, containing 20% of Ham's F 12, 10% of FCS, 100 IU/ml of penicillin, 100 µg/ml of streptomycin and 2.5 µm/ml of amphotericin B.

Once the cells reach confluence, they are recovered with the help of trypsin (1mg/ml) in the presence of EDTA (ethylenediaminetetraacetic acid).

30 Then, they are grown in the "prepared DMEM medium".

35 Before adding the cells in Petri dishes containing a support with a fibrin network, the cells were recovered from the DMEM medium prepared by incubation in a trypsin-EDTA medium (5 times as concentrated) for 5 minutes at 37°C, then 10 ml of a culture medium containing 10% of FCS were added to stop the action of the enzyme. The number of cells in the

medium was determined with the help of a microscope by counting the cells in a Bürker chamber after trypan blue marking. This method will be hereafter named microscope counting method. The resulting number of cells was 5 25,000 cells/ml for a first solution and 125,000 cells/ml for a second solution.

2 ml of the culture medium, containing 50,000 cells and 250,000 cells respectively were added separately in the different Petri dishes respectively 10 containing a lyophilized fibrin network prepared with no added glycerol (Dish 1), a fibrin network prepared with a solution containing about 5% of glycerol (example 4) before lyophilization (Dish 2) and after lyophilization (Dish 3), and a fibrin network prepared with a solution containing about 10% of glycerol (example 5) with 15 lyophilization (Dish 4).

The culture of cells in Petri dishes occurred at 37°C for 2 hours for a first batch of dishes (dishes containing 50,000 cells) and for 11 days for a second 20 batch of dishes (dishes containing 250,000 cells). When the culture time - either 2 hours, or 11 days - expired, the fibrin networks in Petri dishes were fixed by means of a 2.5% glutaraldehyde solution. Figures 15, 16, 17 and 18 are top views of the fibrin layer of the dishes 25 1, 2, 3, and 4 after 2 hours culture, as taken with an electron microscope (Philips XL20 Scanning Electron Microscope). These figures show good cell attachment on fibrin networks in the different dishes, after two hours of culture. The cells are distributed on the upper 30 surface with a regular and flat arrangement.

For the dishes in culture for 11 days at 37°C, a visual examination of dishes was performed during the culture time. This examination showed that, after 8 days of culture, fibrinolysis of the network of 35 dish 1 (fibrin network without glycerol) was visible, whereas no fibrinolysis was perceptible for the networks of dishes 2, 3 and 4 after 8 days of culture.

After 11 days of culture, the number of viable cells was counted for dish 1 and for dishes 2 and 3. The number of viable cells was determined by means of an enzymatic kit, Boehringer Mannheim WST-1 (Catalogue no. 1644807 - batch no. 14890800). The principle of this method is based on the cleavage of a tetrazolium salt, added to the medium, into formazan, by a mitochondrial enzyme (succinate-tetrazolium reductase). This reduction only takes place in viable cells. The formazan color produced by metabolically active cells is quantified by a scanning spectrophotometer (ELISA reader). This determination was made by replacing the culture medium of Petri dishes 1, 2 and 3 by 1 ml of a fresh medium containing 100 μ l of the solution of the WST-1 enzymatic kit. After four hours of incubation at 37°C under an atmosphere containing 7% of CO₂, 100 μ l of the colored solution of each dish were collected for a spectrometer analysis. The difference between the absorbance peak at 450 nm and the absorbance at 655 nm was determined for each solution. The absorbance difference for dishes 2 and 3 was found to be much more important (40 to 50% higher) than for dish 1. The absorbance difference for dishes 1, 2 and 3 was at least four times higher than that of a sample with no cells therein. This analysis proves that the cells in dishes 1, 2 and 3 are viable, and further that the presence of glycerol ensures better cellular viability.

CLAIMS

1. An element provided with a layer based on fibrin- or fibrinogen-containing material, said element comprising (a) a hydrophobic or substantially hydrophobic support, which has a porous part with a thickness of 0.1 to 5 mm, and whose pores, extending across its thickness have a node spacing of 5 to 100 μm , one face of said porous part being treated with a compound based on fibrin and/or a fibrinogen-containing material, and (b) a fibrin-based layer covering said treated surface of the support, characterized in that said fibrin-based layer is substantially uniform and homogeneous on said treated surface, and that the fibrin layer and at least the face of the support in contact with the fibrin layer are substantially free of fibrinogen.

2. An element as claimed in claim 1, characterized in that the fibrin layer and at least a support layer extending across a thickness of 10 μm contains less than 1% by weight, advantageously less than 0.5%, preferably less than 0.1% by weight of fibrinogen which has not reacted, with respect to the weight of the fibrin layer.

3. An element as claimed in claim 1 or 2, characterized in that fibrin extends across the thickness of the treated porous part of the support, from said treated face to a depth of at least 2 μm , both through the pores having an average diameter of 10 to 20 μm and through the pores having an average diameter of more than 20 μm .

4. An element as claimed in any preceding claim, wherein the porous part of the support has a substantially homogeneous and uniform porosity over the treated face, characterized in that some fibrin extends homogeneously and uniformly across the thickness of the porous part of the support to a depth of at least 10 μm .

5. An element as claimed in claim 2, characterized in that the porous support contains fibrinogen in a layer which is at a distance of more than 10 μm from the face in contact with the fibrin layer.

6. An element as claimed in claim 5, characterized in that fibrinogen extends across the thickness of the support to a depth of at least 20 μm .

7. An element as claimed in any preceding claim, characterized in that at least the face of the fibrin layer opposite to the one contacting the porous support is stabilized.

8. An element as claimed in claim 7, characterized in that said fibrin-based layer is at least partially cross-linked, to form a network of adjacent alveoli.

9. An element as claimed in any preceding claim, characterized in that said layer is provided with cells and/or proteins, particularly with proteins mediating cell-fibrin bonds.

10. An element as claimed in any preceding claim, characterized in that the cross-linked fibrin-based layer which covers the porous part of the support, when measured in the dry state, is 0.5 to 100 μm thick, preferably 2.5 to 50 μm thick, with alveoli being formed between the cross-linked fibrin-based molecules or bonds, or fibers, said alveoli having a volume of 5 to 25 μm^3 , the average thickness or height of said alveoli being of 1 to 5 μm , particularly of 1 to 3 μm .

11. An element as claimed in any preceding claim, characterized in that the pores of the support part, covered by said fibrin layer have inner faces at least partially covered by a water-soluble or substantially water-soluble protein.

12. An element as claimed in claim 11, characterized in that the support face opposite to the

treated face is at least partially covered by a water-soluble or substantially water-soluble protein.

13. An element as claimed in any preceding claim, characterized in that at least the pores of the 5 porous part of the support are covered by a water-soluble or miscible polar organic additive.

14. An element as claimed in claim 13, characterized in that the network of cross-linked fibrin fibers is at least partially covered by and/or contains 10 a water-soluble or miscible polar additive, preferably an additive selected in the group comprising glycerol, sugars and mixtures thereof.

15. An element as claimed in any preceding claim, characterized in that it has a moisture content of less than 0.5% by weight, preferably of less than 0.1% by weight.

16. An element as claimed in any preceding claim, characterized in that fibronectin is attached to the fibrin layer, the fibronectin content, as compared 20 to the fibrin and fibronectin weight in the layer being of 0.5 to 15%.

17. An element as claimed in any preceding claim, characterized in that the fibrin layer contains calcium in the order of 1 to 100 µg, preferably of 1 to 25 50 µg of calcium per cm³ of the fibrin layer volume.

18. An element as claimed in claim 14, characterized in that calcium takes the form of calcium chloride.

19. An element as claimed in any preceding 30 claim, characterized in that the support has two superposed fibrin layers, the layer in contact with the support having alveoli with larger volumes as compared with the alveoli of the layer which covers the fibrin layer in contact with the support.

20. An element as claimed in any preceding 35 claim, characterized in that the support is biocompatible and/or biodegradable.

21. A process for preparing an element as claimed in any claim 1 to 20, wherein at least one porous part of a first face of a porous support is placed in contact with an aqueous solution containing fibrin or a fibrinogen-containing material, wherein the face of the porous part of the support opposite to said first face is homogeneously and uniformly submitted to a suction force to suck the solution, at least partly, through said porous part, thus ensuring the deposition of a layer based on fibrin or on fibrinogen-containing materials, homogeneously and uniformly with respect to said porous part, and the diffusion of at least the solution water through the porous part of the porous support as well as the penetration of fibrin or fibrinogen through the porous support.

22. A process as claimed in claim 21, wherein the face of the support opposite to said first face, is submitted to a pressure of less than $0.8 \cdot 10^5$ Pa, and wherein a pressure difference is created between the two faces of the porous part of at least $0.3 \cdot 10^5$ Pa.

23. A process as claimed in claim 22, characterized in that the support face opposite to said first face is submitted to a pressure of less than $0.5 \cdot 10^5$, preferably less than or equal to $0.4 \cdot 10^5$ Pa.

24. A process as claimed in claim 22 or 23, characterized in that the support face opposite to the first face is intermittently submitted to a first pressure, of less than $0.8 \cdot 10^5$ Pa, preferably less than $0.4 \cdot 10^5$ Pa, and to a second pressure of less than $0.8 \cdot 10^5$ Pa, preferably less than $0.4 \cdot 10^5$ Pa, the first pressure being at least 5% higher than the second pressure.

25. A process as claimed in any claim 22 to 24, wherein the face of the porous support opposite to said first face is submitted to a pressure of less than $0.8 \cdot 10^5$ Pa, and exposed to a temperature of 0 to 100°C, preferably to a temperature of 15 to 60°C.

26. A process as claimed in claim 21, wherein the face of the porous support opposite to said first face is submitted to a solution selected so as to create a reverse osmosis, causing the diffusion of at least the solution water in contact with the first face through the porous part of the support.

27. A process as claimed in any claim 21 to 26, characterized in that a solution is used which contains 5 to 20 mg/ml of fibrinogen-containing materials.

28. A process as claimed in claim 27, characterized in that a solution is used which contains 5 to 20 mg/ml of fibrinogen-containing materials and 0.01 to 10 units of thrombin per ml.

29. A process as claimed in claim 28, characterized in that a solution is used which contains 5 to 20 mg/ml of fibrinogen-containing materials, factor XIII, and 0.01 to 2 units of thrombin per ml.

30. A process as claimed in claim 29, characterized in that a solution is used which contains 0.1 to 10 units of factor XIII per ml.

31. A process as claimed in any claim 27 to 30, characterized in that the solution contains 1 to 40 millimoles of calcium chloride per ml.

32. A process as claimed in any claim 27 to 31, characterized in that the solution contains 0 to 20% by weight, advantageously 3 to 15%, preferably 5 to 10% of a water-soluble or miscible polar organic additive.

33. A process as claimed in claim 32, characterized in that the additive is glycerol.

34. A process as claimed in any claim 21 to 33, characterized in that, during a first step, at least one portion of a first face of a porous support is placed in contact with a solution containing fibrin and/or fibrinogen-containing materials, wherein the face of the porous support opposite to said first face is submitted to a suction force, thus ensuring a diffusion

of at least the solution water across the thickness of the porous support and a penetration of fibrin or fibrinogen across the thickness of the porous support, homogeneously and uniformly with respect to said porous part of the first face and in that, during a second step, the fibrin and/or fibrinogen layer is stabilized.

5 35. A process as claimed in any claim 21 to 34, characterized in that a contact is provided between said part of the first face and a moving aqueous 10 solution.

15 36. A process as claimed in any claim 21 to 35, characterized in that an aqueous solution is used which contains a wetting agent to fill the pores of the porous support before placing said support in contact with the solution containing fibrin or fibrinogen-containing materials.

20 37. A process as claimed in any claim 21 to 36, characterized in that the fibrin layer is submitted to a drying step, possibly preceded by a washing step.

25 38. A process as claimed in claim 37, characterized in that this drying operation is effected at least partially by lyophilization, advantageously at a temperature of -30°C and -100°C, preferably at a temperature of -40°C to -70°C.

30 39. A process as claimed in any claim 28 to 38, characterized in that at least for a part of the deposit of the layer based on fibrin or on fibrinogen-containing materials, the concentration of fibrin or fibrinogen-containing materials in the solution in contact with the first face is controlled in order to ensure a substantially constant water diffusion through the support.

35 40. A process as claimed in any claim 21 to 39, characterized in that a biocompatible and/or biodegradable porous support is used.

41. A process as claimed in any claim 21 to 40, characterized in that the porous part is treated

with an aqueous solution which advantageously contains a wetting agent, a protein or a polar organic additive, or a mixture thereof, before bringing the solution containing fibrin and/or fibrinogen-containing materials 5 in contact with said porous part.

42. A filter including a membrane consisting of an element as claimed in any claim 1 to 20.

10 43. A bioreactor including a membrane consisting of an element as claimed in any claim 1 to 20.

44. An implant consisting of an element as claimed in any claim 1 to 20.

15 45. An artificial skin produced from an element as claimed in any claim 1 to 20.

FIG 1

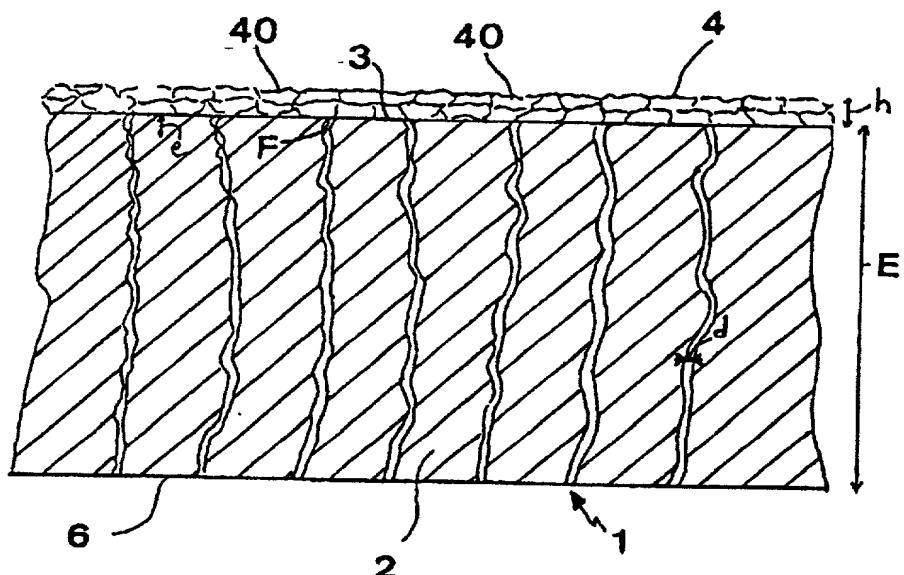


FIG 2

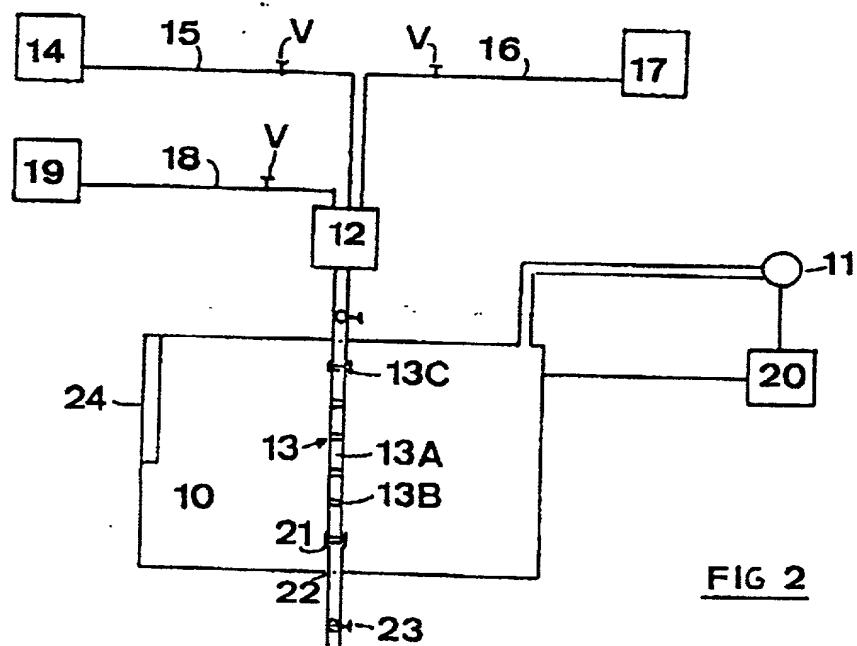


Fig. 3

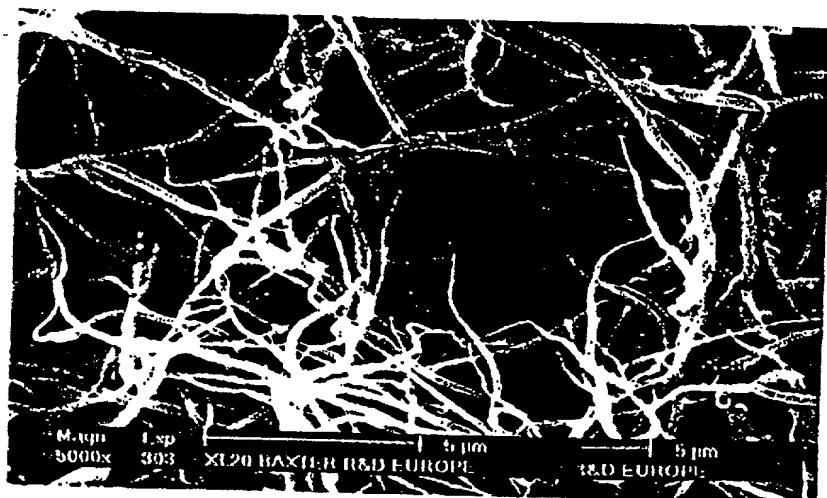


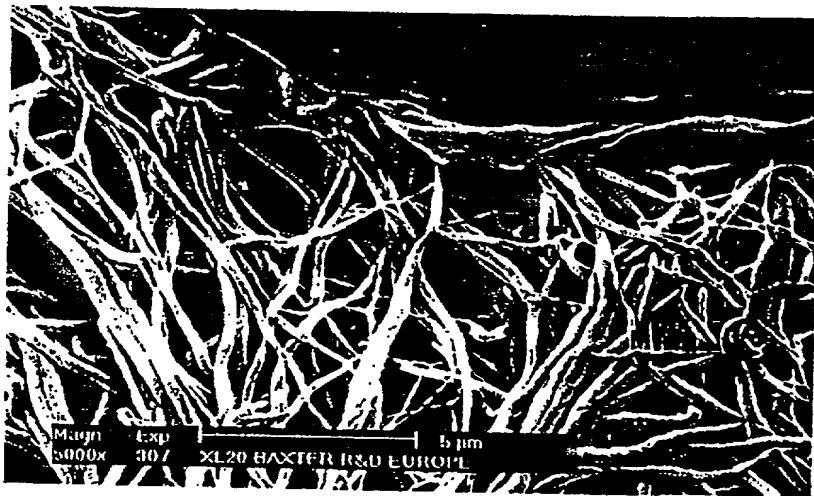
Fig. 4



Fig.5



Fig.6



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Fig. 7



Fig. 8

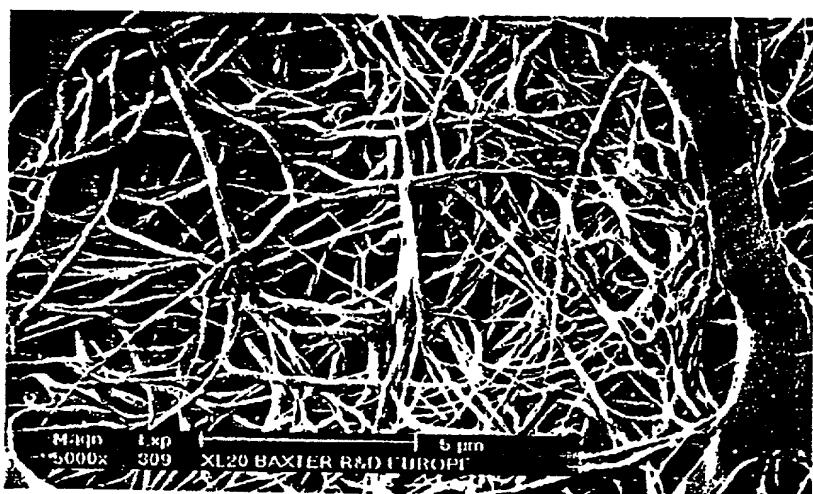


Fig.9.

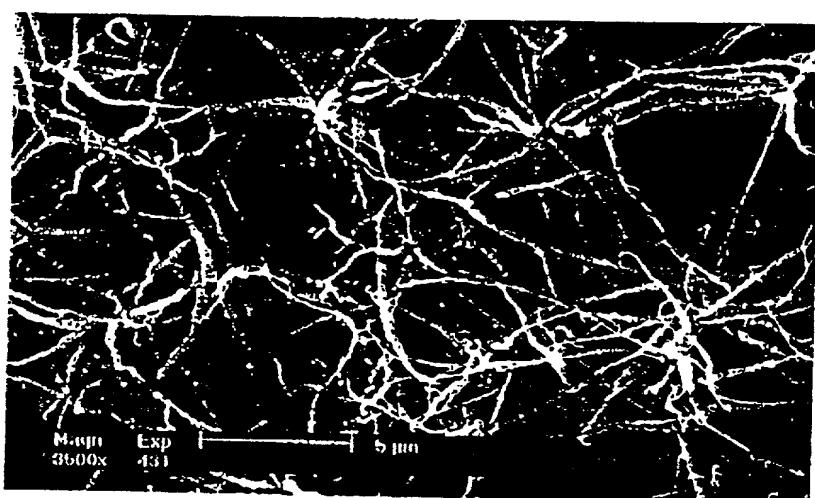


Fig. 10

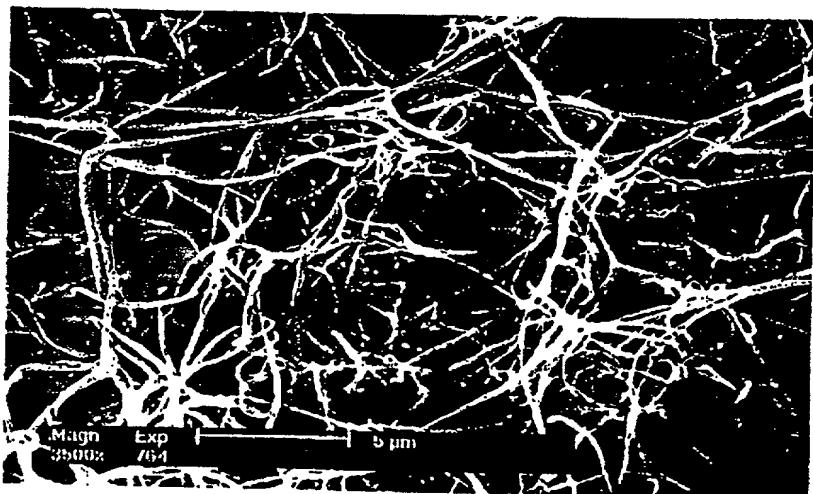


Fig.11

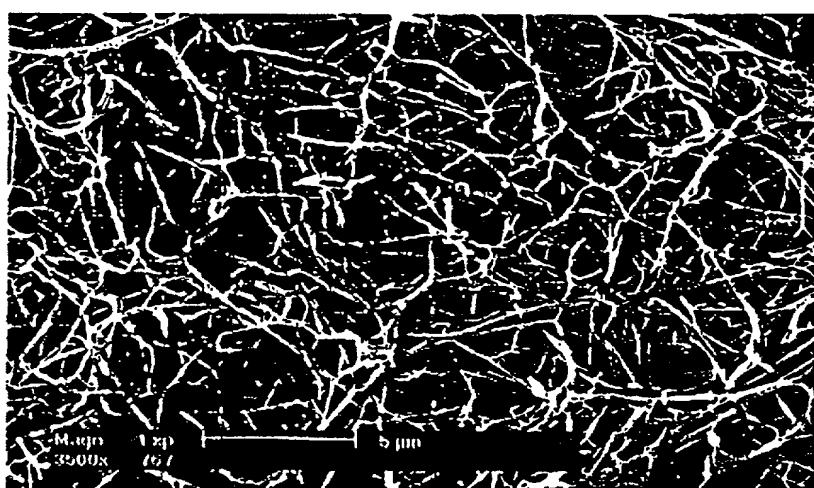


Fig. 12



Fig.13

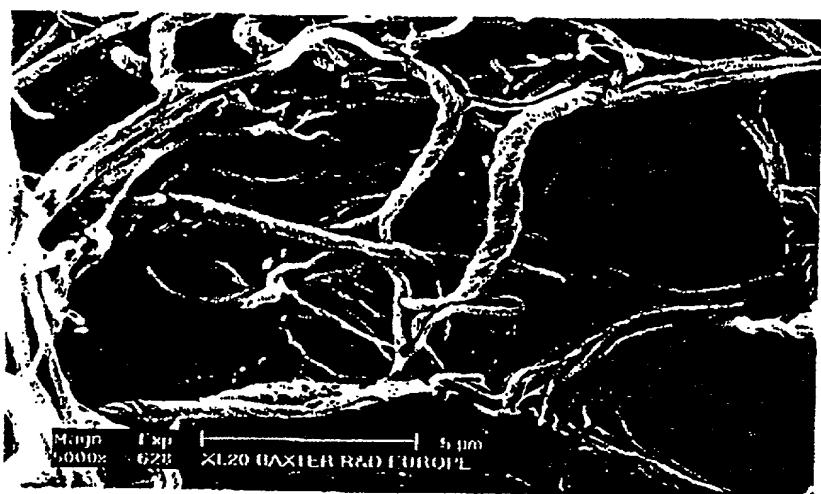


Fig. 14



Fig.15

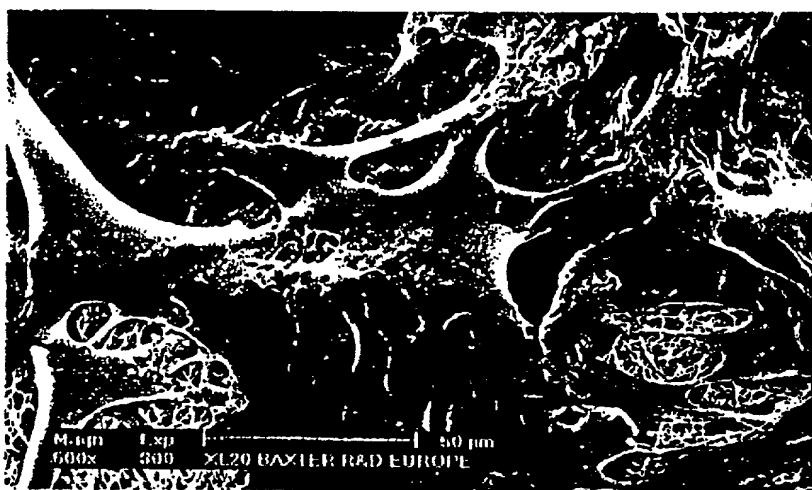


Fig. 16



Fig.17

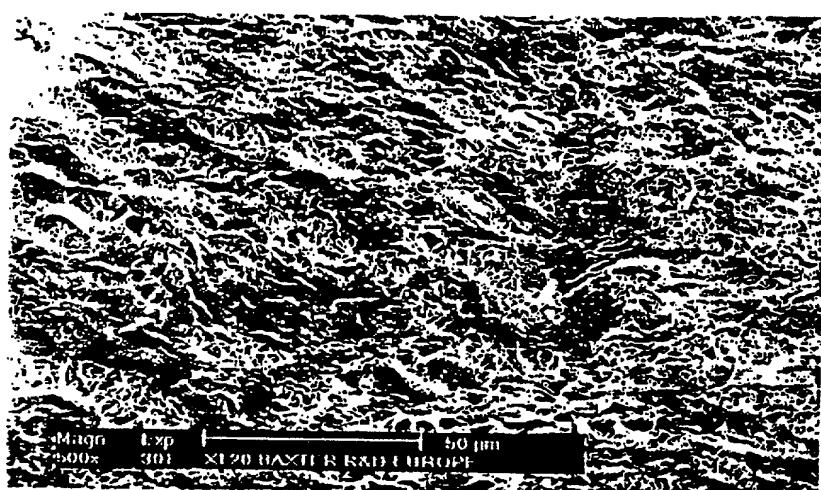
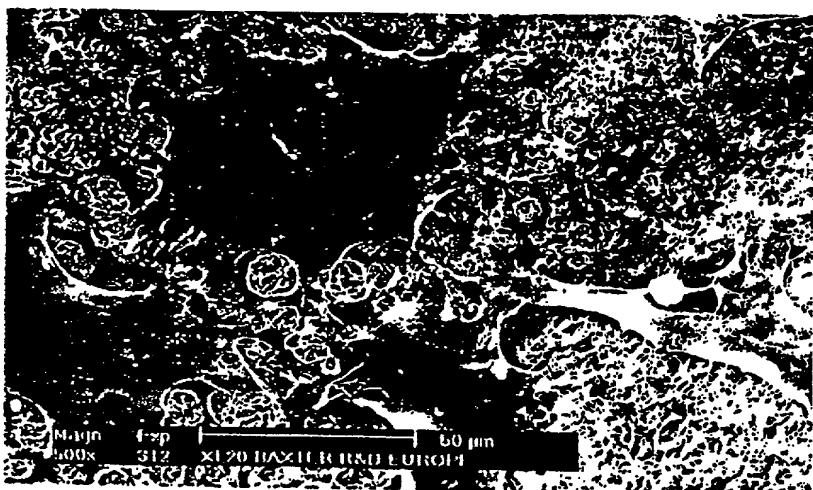


Fig. 18



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Fig. 19

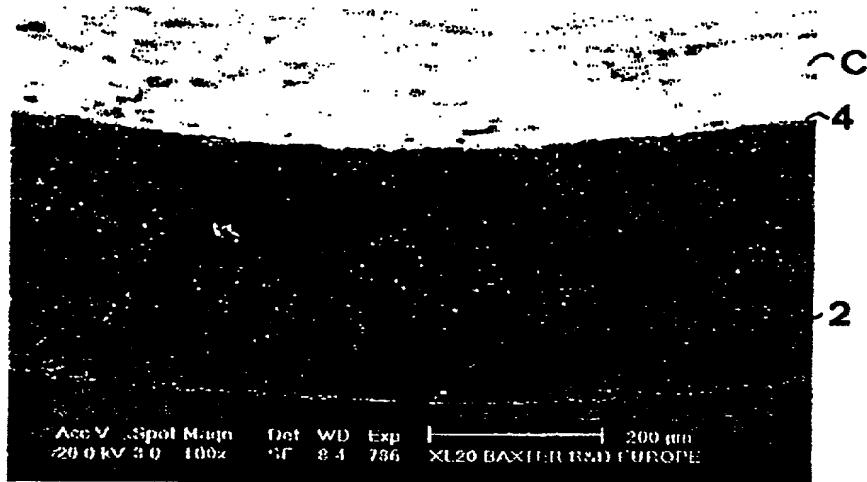
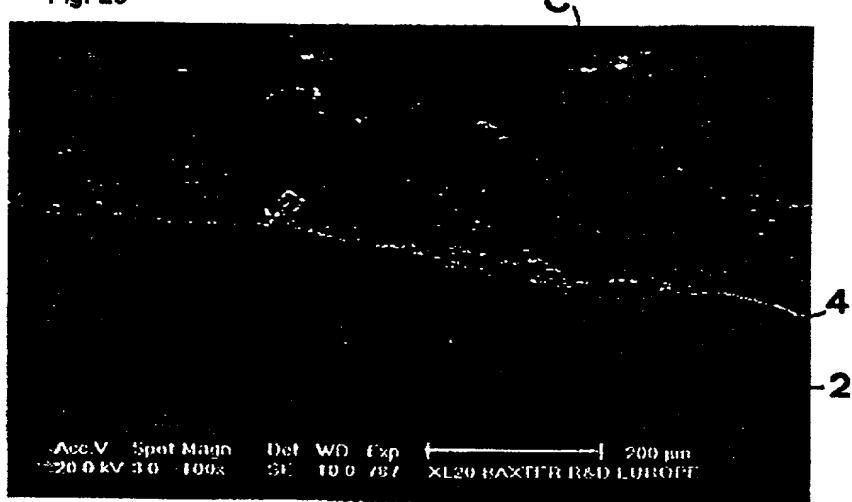


Fig. 20



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Fig.21

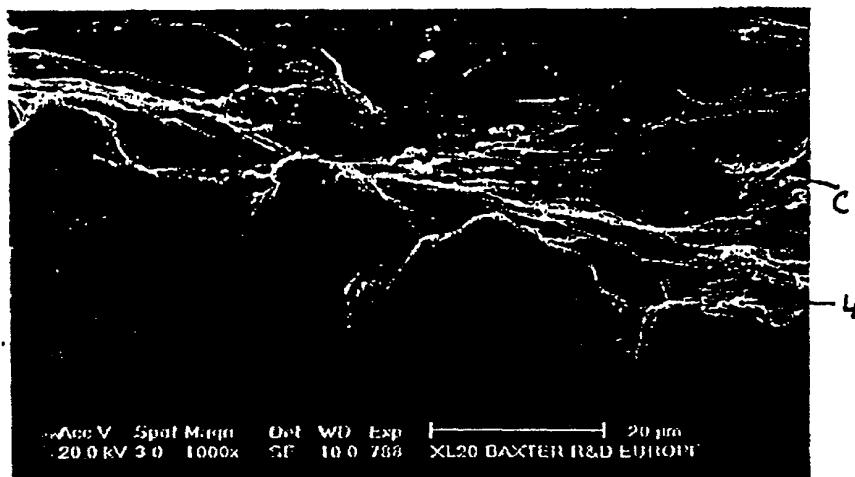
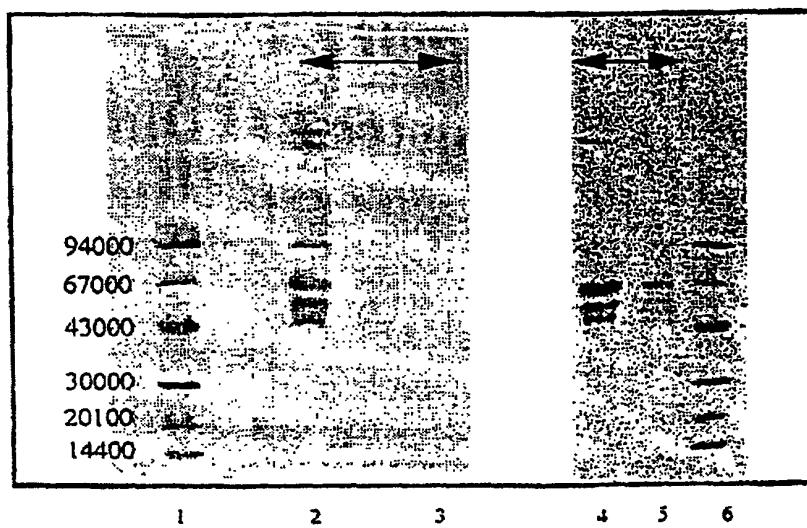


FIG 22



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/121736

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

DECLARATION AND POWER OF ATTORNEY

As below-named inventors, we hereby declare that:

Our residences, post office addresses and citizenships are as stated below next to our names.

We believe we are the original, first and joint inventors of the subject matter which is claimed and for which a patent is sought on the invention entitled "**ELEMENT PROVIDED WITH A FIBRIN LAYER, PREPARATION AND USE THEREOF**", filed on May 4, 2001, as Application No. _____.

We hereby state that we have reviewed and understand the contents of the above-identified Specification, including the Claims, as amended by any Amendment referred to above.

We acknowledge the duty to disclose information which is material to patentability as defined in 37 C.F.R. § 1.56.

We hereby claim foreign priority benefits under 35 U.S.C. § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT international application which designated at least one country other than the United States, listed below and have also identified below any foreign application for patent or inventor's certificate, or PCT international application having a filing date before that of the Application on which priority is claimed:

Prior Foreign Application(s):

		Priority <u>Claimed</u>
9800796	Belgium	X
Number	Country	Yes No
	4 November, 1998	
	Day/Month/Year	
	Filed	

We hereby claim the benefit under 35 U.S.C. § 119(e) of any United States provisional application(s) listed below:

(None)
Application Number

Filing Date

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We hereby claim the benefit under 35 U.S.C. § 120 of any United States Application(s), or § 365(c) of any PCT International Application designating the United States, listed below and, insofar as the subject matter of each of the Claims of this Application is not disclosed in the prior United States or PCT International Application in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 C.F.R. § 1.56 which became available between the filing date of the prior Application and the national or PCT international filing date of this Application:

<u>PCT/US99/25955</u>	<u>4 November, 1999</u>	<u>Pending</u>
Application Number	Filing Date	Status

We hereby appoint the following attorney(s) and/or agent(s) to prosecute this Application and transact all business in the Patent and Trademark Office connected therewith.

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Joseph B. Barrett	<u>34,769</u>	Roger H. Stein	<u>31,882</u>	Edward L. Bishop	<u>39,110</u>
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		Bradley F. Rademaker	<u>35,331</u>	Brent A. Hawkins	<u>44,146</u>
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We hereby declare all statements made herein of our own knowledge are true and all statements made on information and belief are believed to be true; and further, that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States

Declaration and Power of Attorney

Baxter Reference No. CRT-5463

W&W File No. 1417S P 585

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Code and that such willful false statements may jeopardize the validity of the Application or any patent issued thereon.

1.0 Full Name of First Joint Inventor: Yves Delmotte

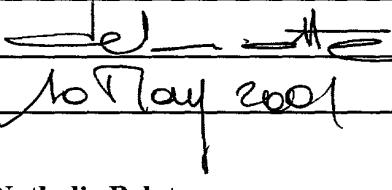
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Inventor's Signature: 

Date: 10 May 2001

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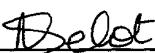
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Citizenship: Belgium

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Inventor's Signature: 

Date: 10 May 2001

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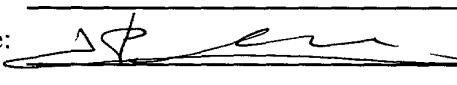
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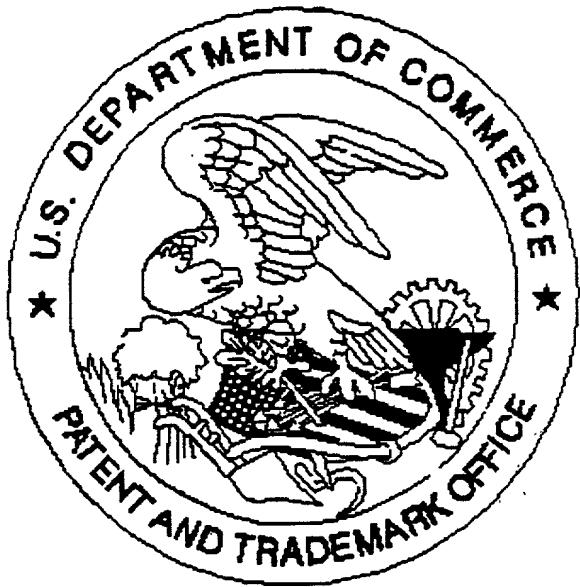
Citizenship: Belgium

Mailing Address: Same as above.

Inventor's Signature: Nicole Tasiaux

Date: 10 May 2001

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